SEARCHER QUEST FORM

Scientific and Ecchnical Information Center

OCT -4. 2002

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Requester's Full Name: Leigh	h Maier	Evaminar #	77012	STIE)
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Mail Box and Bldg/Room Locati	on: 7403	Results Format Pre	ferred (circle): (PA	DED DICK E MAII
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If more than one search is sub	mitted, please prior	ritize searches in	order of need.	
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utility of the invention. Define any term known. Please attach a copy of the cove	us uiai may nave a snecia	l meaning Citye ever	nples or relevant citat	tions, authors, etc, if
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Title of Invention:	Bib she	eet attach	ed	0.0
Inventors_(please provide full names):			<u> </u>	
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Earliest Priority Filing Date:	V	*.*		2
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	*			-30XL44QQ -
			jan.delaval@us	pro.gov .
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TAFF.USE ONLY	Type of Search	Vendor	s and cost where app	olicable
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WWW/Internet

Other (specify)

PTO-1590 (8-01)

Online Time:

Date Completed:

Searcher Prep & Review Time

Date Searcher Picked Up: (3) 10) 02

10/10/02

Litigation

Fulltext Patent Family

Other

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STRUCTURE FILE UPDATES: 9 OCT 2002 HIGHEST RN 460312-12-3 DICTIONARY FILE UPDATES: 9 OCT 2002 HIGHEST RN 460312-12-3

TSCA INFORMATION NOW CURRENT THROUGH MAY 20, 2002

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

=> d ide can l1

- L1ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS RN 11138-66-2 REGISTRY CN Xanthan gum (9CI) (CA INDEX NAME) OTHER NAMES: Actigum CX 9 CN
- CN ADM 40
- CN B 1459
- Biopolymer 9702 CN CN
- Biopolymer XB 23
- CNBiozan R
- Bisfect XA 200 CN
- CN Bistop .
- CN Chemicogel
- Echogum CN
- Echogum F CN
- Echogum RD CN
- Echogum SF CN
- CN Echogum T
- CN Ekogum
- Ekogum ketorol CN
- Enorflo X CN
- CN Flocon 1035
- Flocon 4800 CN
- CN Flocon 4800C
- CN Flodrill S
- CN Galaxy XB
- Gums, xanthomonas CN
- Idvis CN
- CN Jungbunzlauer ST
- K 5C151 CN
- K 9C57 CN
- Kelco BT CN
- CN Kelflo
- Keltrol CN
- Keltrol CG CN
- Keltrol F CN
- CN Keltrol RD

Jan Delaval Reference Librarian Biotechnology & Chemical Library CM1 1E07 - 703-308-4498 jan.delaval@uspto.gov

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CN
     Keltrol SF
     Keltrol T
CN
     Keltrol TF
CN
     Keltrol TF 1000
CN
     Kelzan
CN
CN
     Kelzan 140X
     Kelzan AR
CN
     Kelzan ASX
CN
     Kelzan D
CN
     Kelzan F
CN
     Kelzan M
CN
     Kelzan MF
CN
     Kelzan S
CN
     Kelzan SS 4000
CN
     Kelzan T
CN
     Kelzan XC
CN
     Kelzan XCD
CN
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
     12673-42-6, 12771-06-1, 9088-32-8, 54511-23-8, 56592-13-3, 98112-77-7,
DR
     51811-95-1, 37189-49-4, 37279-85-9, 37332-19-7, 37383-52-1, 80450-59-5,
     85568-76-9, 82600-55-3, 39393-27-6, 39444-54-7
     Unspecified
MF
     PMS, COM, MAN
CI
    Manual registration, Polyester, Polyester formed
PCT
LC
     STN Files: AGRICOLA, ANABSTR, AQUIRE, BIOBUSINESS, BIOSIS, BIOTECHNO,
       CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CHEMLIST, CIN,
       CSCHEM, DDFU, DRUGU, EMBASE, ENCOMPLIT, ENCOMPLIT2, ENCOMPPAT,
       ENCOMPPAT2, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*, MSDS-OHS,
       NAPRALERT, NIOSHTIC, PIRA, PROMT, TOXCENTER, TULSA, USPAT2, USPATFULL,
         (*File contains numerically searchable property data)
     Other Sources: DSL**, EINECS**, TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
            6852 REFERENCES IN FILE CA (1962 TO DATE)
             236 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
            6886 REFERENCES IN FILE CAPLUS (1962 TO DATE)
            1: 137:222159
REFERENCE
REFERENCE
            2:
                137:222095
REFERENCE
                137:222032
            3:
REFERENCE
                137:221941
            4:
REFERENCE
            5:
                137:221810
REFERENCE
            6:
                137:221757
REFERENCE
            7:
                137:218774
REFERENCE
            8:
                137:216182
REFERENCE
            9:
                137:212293
REFERENCE 10: 137:208425
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(FILE 'HOME' ENTERED AT 07:45:35 ON 10 OCT 2002) SET COST OFF

FILE 'REGISTRY' ENTERED AT 07:46:08 ON 10 OCT 2002 E XANTHAN GUM/CN

T 1		1		XANTHAN GUM/CN
L1		T	5	E3
	FILE	' HCAPI		S' ENTERED AT 07:46:23 ON 10 OCT 2002
L2		6898		
L3	•			L1 (L) DEACET?
L4				L3 AND L2
L5				XANTHAN (A) GUM
כת		0031		
				DEACETYLATION/CT
T. C		1000		E3+ALL
L6		1069		
				E4+ALL
L7		597		
				E9+ALL
rs		987		E4, E3+NT
			Ε	E10+ALL
			E	E8+ALL
L9		1466	S	E2, E4
L10		14	S	L2,L5 AND L6-L9
L11		43	S	L2, L5 AND DEACET?
L12		13	S	L2, L5 AND DEACYL?
L13				L4,L10-L12 AND L2-L12
				LANGLOIS B/AU
L14		67		E3-E5, E11-E13
L15				L14 AND L13
L16				L14 AND L2, L5
што		-		RHODIA/PA, CS
L17		1137		E3, E4
L18				L2,L5 AND L17
L19				L13 AND L15, L16, L18
L20		31		L18 NOT L19
				DRILLING FLUID/CT
- 01		2554		E4+ALL
L21		1554		E2, E3, E1+NT
				E8+ALL
L22		1151		E2, E1+NT
				E7+ALL
L23		1561		E3+NT
		•		DRILLING FLUID/CT
				E4+ALL
			Ε	E9+ALL
L24		2374		E4, E3+NT
			Ε	E13+ALL
L25		5086	S	E4, E3+NT
			Ε	E12+ALL
L26		2374	S	E4, E3+NT
			Ε	E2+ALL
L27		12718	S	E3, E2+NT
				DRILLING FLUIDS/CT
L28		402		
			Ε	E3+ALL
				E11+ALL
L29		388		
L30				L2, L5 AND L21-L29
L31				L30 AND L20
L32				L30 AND L13
L33				L31, L32
L34				
				L33 AND LANGLOIS ?/AU
L35		56	5	L13 NOT L34

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L36
              0 S L35 AND L30
L37
              1 S L35 AND FUEL?/SC, SX
L38
              2 S L34, L37
L39
             55 S L35 NOT L38
L40
             55 S L39 AND XANTHAN
L41
             47 S L40 AND GUM
             8 S L40 NOT L41
L42
L43
             55 S L39-L42
L44
             49 S L43 AND (PD<=20000114 OR PRD<=20000114 OR AD<=20000114)
                SEL DN AN 3 4 14 25 30 31 35 44 45 46 47
             11 S L44 AND E1-E33
L45
L46
             13 S L38, L45
L47
             44 S L43 NOT L46
                SEL DN AN 5 6 11 16 17 18 19 20 25 26 28 31 32 33 34 35 38 39 4
L48
           21 S L47 AND E34-E96
L49
             34 S L46, L48
L50
              9 S L2, L5 AND (DE ACET? OR DE ACYL? OR NONACET? OR NONACYL? OR NO
L51
              4 S L50 AND L49
L52
              5 S L50 NOT L51
L53
             4 S L52 NOT SUBSTRATE/TI
L54
             38 S L49, L51, L53
           8536 S L2, L5 OR XANTHAN
L55
L56
             3 S L55 AND ?PENTAMER?
L57
             40 S L54, L56
L58
             40 S L57 AND L2-L57
L59
             9 S L58 AND ?ACYL?
             36 S L58 AND ?ACETYL?
L60
             40 S L58-L60
L61
            161 S L55 AND C09K007/IC, ICM, ICS
L62
L63
              1 S L62 AND L61
L64
             40 S L61, L63
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FILE 'REGISTRY' ENTERED AT 08:21:12 ON 10 OCT 2002

=> fil hcaplus

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FILE COVERS 1907 - 10 Oct 2002 VOL 137 ISS 15 FILE LAST UPDATED: 9 Oct 2002 (20021009/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

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- L64 ANSWER 1 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 2002:605427 HCAPLUS
- TI Novel application of synergistic guar/non-acetylated xanthan gum mixtures in hydraulic fracturing
- AU Fischer, C. C.; Navarrete, R. C.; Constien, V. G.; Coffey, M. D.; Asadi, M.
- CS Constien & Associates, USA
- SO SPE International Symposium on Oilfield Chemistry, Conference Proceedings, Houston, TX, United States, Feb. 13-16, 2001 (2001), 485-496 Publisher: Society of Petroleum Engineers, Richardson, Tex. CODEN: 69CZJT
- DT Conference; (computer optical disk)
- LA English
- AB Fracturing fluids have traditionally been viscosified with guar and guar derivs. Non-acetylated xanthan is a variant of xanthan gum which when combined with guar in soln. develops a synergistic interaction that generates superior viscosity and particle transport at low polymer concns. These water-base linear fluids

particle transport at low polymer concns. These water-base linear fluids have improved low shear viscosity at concns. at or below 25 lb/1,000gal when compared to fluids viscosified using a single viscosifier such as guar or xanthan gum. The polymer mixts. can be

crosslinked to provide enhanced viscosity at higher temps.

RETABLE

Referenced Author (RAU)	Year VOL (RPY) (RVL	•	Referenced Work Referenced (RWK) File
Anon Dea, I Gulbis, J Harris, P Mwamufiya, I Nimerick, K Pope, D Roodhart, L Samuel, M Shah, S Talashek, T Tehrani, A Unwin, A Willberg, D	1998 1993	-+	Recommended Practice Industrial Gums Reservoir Stimulatio Paper SPE 38621, pre PhD Thesis, Princeto Paper SPE 35638, pre Paper SPE 31094, pre Paper SPE 13905, pre Paper SPE 38622, pre Paper SPE 49040, pre Personal Communicati J Rheol Paper SPE 29649, pre Paper SPE 38620, pre

- L64 ANSWER 2 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 2001:163308 HCAPLUS
- DN 134:349490
- TI Effect of a range of microbial polysaccharides on the diffusion of manganese ions using spatially resolved NMR relaxometry
- AU Hart, T. D.; Hill, R. J.; Glover, P. M.; Lynch, J. M.; Chamberlain, A. H. L.
- CS School of Biological Sciences, University of Surrey, Guildford, GU2 5XH, UK
- SO Enzyme and Microbial Technology (2001), 28(4-5), 370-375 CODEN: EMTED2; ISSN: 0141-0229
- PB Elsevier Science Ireland Ltd.
- DT Journal
- LA English
- AB In accordance with the theory of contact exchange, it is hypothesized that the presence of neg. charge in microbial exopolysaccharides increases the rate of cation transport. These typically acidic materials may provide a fast-track for the diffusion of nutrient cations through the polymer layer for uptake at the organism cell surface. We have measured the diffusion coeff. of a model cation, Mn2+, through xanthan, deacetylated xanthan, scleroglucan and chitosan using

spatially resolved NMR relaxometry. The concn. of Mn2+ in soln. was measured by recording the change in the spin-spin (T2) relaxation time of water 1H over time in compartments either side of a polymer layer. This approach provides a sensitive, in situ, non-invasive method of measuring the rate of diffusion of paramagnetic cations through hydrophilic polysaccharides. The neg.-charged polysaccharides, xanthan and de-acetylated xanthan, permitted a

significantly faster rate (2-2.5.times.) of cation transport compared to the uncharged polymer, scleroglucan. The pos.-charged polysaccharide chitosan reduced the rate of Mn2+ diffusion to around half the value obtained for scleroglucan. These results suggest that the presence and nature of fixed charges on the polysaccharide mol. affects the rate of cation transport in accordance with the theory of contact exchange. The presence of neg. charge on microbial exopolysaccharides may thus improve the availability of nutrient cations at the organism cell surface.

IT 11138-66-2, Xanthan 11138-66-2D,

Xanthan, deacetylated

RL: PEP (Physical, engineering or chemical process); PROC (Process) (effect of a range of microbial polysaccharides on the diffusion of manganese ions using spatially resolved NMR relaxometry)

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Referenced Author (RAU)	(RPY)	(RVL)	PG (RPG)	(RWK)	Referenced File
Bloembergen, N	1961		1842		HCAPLUS
Corpe, W	1980	İ	1105	Adsorption of microo	HCAPLUS
Efron, B	1977	7	1	Ann Stat	ĺ
Foster, R	1983	İ	Ì	Ultrastructure of th	ĺ
Geddie, J	11993	174	467	J Appl Bacteriol	HCAPLUS
Geesey, G	1989	1	325	Metal ions and bacte	
Glauser, R	1960	5	1	Agrochimica	HCAPLUS
Harned, H	1958	1	1	The physical chemist	1
Hart, T	12000	1	1	Environ Microbiol, i	1
Hart, T	1999	24	1339	Enzyme Microb Techno	HCAPLUS
Hassler, R	1990	6	1182	Biotechnol Prog	HCAPLUS
Holzwarth, G	1985	126	271	Dev Ind Microbiol	HCAPLUS
Jeanes, A	1961	5	519	J Appl Polymer Sci	HCAPLUS
Jenny, H	1961	5	281	Agrochimica	1 .
Jenny, H	1961	1	1665	Growth in living sys	1
Jenny, H	1966	25	1265	Pl Soil	HCAPLUS
La Paglia, C	1997	63	3158	Appl Environ Microbi	HCAPLUS
Luz, Z	1965	43	13750	J Chem Phys	HCAPLUS
Meares, P	1958	55	1273	J Chim Phys	HCAPLUS
Nambier, G	1976	4 4	1267	Pl Soil	1
Patil, S	1993	173	153	J Radional Nuclear C	HCAPLUS
Ramamoorthy, S	1977	166	527	J Theor Biol	HCAPLUS
Sollner, K	11974	153	1267	J Dental Res	HCAPLUS
Tako, M		489	268	ACS Symposium Series	HCAPLUS

- L64 ANSWER 3 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 2001:94303 HCAPLUS
- DN 134:310527
- TI Enhanced compatibility of xanthan variants in phosphate systems
- AU Swazey, John
- CS CP Kelco, UK
- SO Research Disclosure (2001), 441(Jan.), P5-P8 (No. 441005) CODEN: RSDSBB; ISSN: 0374-4353
- PB Kenneth Mason Publications Ltd.
- DT Journal; Patent
- LA English

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

PI RD 441005 20010110

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PRAI RD 2001-441005 20010110
     Nonacetylated xanthan (NAX), nonpyruvylated
     xanthan (NPX), and nonacetylated, nonpyruvylated
     xanthan (NPNAX) were tested in ammonium polyphosphate soln.
     (10-34-0) and diammonium phosphate. Hydration is typically not possible
     in fluids contg. >45% 10-34-0 soln. with xanthan products;
     however, with NAX hydration was possible in solns. contq. .gtoreq.60%
     10-34-0. For NPNAX hydration was possible in solns. contg. .gtoreq.70%
     10-34-0. Xanthan gum variants also performed well in
     diammonium phosphate. Compatibility of prehydrated xanthan
     gum variants with phosphates was tested also. Results indicated
     improved compatibility of xanthan variants in systems with high
     phosphate levels. Findings extend xanthan functionality to
     formulations where std. xanthan gum is incompatible or
     unstable.
ΙT
     11138-66-2D, Xanthan gum, deacetylated
     and (or) depyruvylated
     RL: PEP (Physical, engineering or chemical process); PRP (Properties);
     PROC (Process)
        (enhanced compatibility of xanthan variants in phosphate
        systems)
L64
    ANSWER 4 OF 40 HCAPLUS COPYRIGHT 2002 ACS
ΑN
     2000:768961 HCAPLUS
DN
     133:295710
     Viscosity-stable low-acetylated xanthan gum
TΙ
     for food use
IN
     Zablocki, Linda J.; Bousman, W. Scott; Solanki, Yogesh; Milovanovic, Susan
     B.; King, Alan
PA
     Monsanto Company, USA
SO
     U.S., 9 pp.
     CODEN: USXXAM
DT
     Patent
LA
     English
FAN.CNT 1
                      KIND
                           DATE
                                           APPLICATION NO.
     PATENT NO.
                                                            DATE
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                            20001031
                                           US 1999-347259
                                                            19990706 <--
                            20010111
     WO 2001001793
                      A1
                                           WO 2000-US17764 20000629 <--
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR,
             CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU,
             ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
             LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE,
             SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW,
             AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
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                           20020529
                                           EP 2000-941759
                                                           20000629 <--
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL
PRAI US 1999-347259
                      Α
                            19990706
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     WO 2000-US17764
                            20000629
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AΒ
    An acidic edible liq. compn. comprises a low-acetylated
     xanthan gum in an amt. effective to sustain the initial
     viscosity of the compn. for at least about four months.
                                                              Preferred compns.
     include beverages and syrups. Beverages include carbonated and
     non-carbonated soft drinks, still beverages, fruit-juice-type beverages,
     squashes and cordials, alc. and nonalcoholic, their concs. and mixts.
     thereof. A method for stabilizing an acidic edible liq. compn. comprises
     admixing an effective amt. of a low-acetylated xanthan
     gum to maintain the initial viscosity of the compn. for at least
```

about four months under typical storage conditions. Thus,

deacetylated xanthan gum (1.0% wt./wt.;

acetate content 1%) and sodium benzoate (0.33% wt./wt.) were mixed with deionized water; 150 g of the compn. was dild. to 500 g by the addn. of deionized water. The compn. was titrated to pH 4.0 with phosphoric acid. The initial viscosity was 195 cP. After 39 wk storage at ambient temp., e.g., 22.degree., the pH was 3.9 and the viscosity was 177.5 cp; the loss in viscosity during storage was 17.5 cP or about 9%.

IT 11138-66-2, Xanthan gum

RL: FFD (Food or feed use); PRP (Properties); BIOL (Biological study); USES (Uses)

(deacetylated; viscosity-stable low-acetylated
xanthan gum for food use)

RETABLE

Referenced Author (RAU)	Year (RPY)			Referenced Work (RWK)	Referenced File
	- •	+====· '	+======	-+====================================	· • • • • • • • • • • • • • • • • • • •
Anon	1997	I	1	WO 9746656	HCAPLUS
Campaigne	1979		[US 4154654	HCAPLUS.
Cheetham, N	1985	5	399	Carbohydr Polym	HCAPLUS
Doherty	1996	[1	US 5514791	HCAPLUS
Hassler, R	1990	6	182	Biotechnol Prog	HCAPLUS
Jansson, P	1975	45	275	Carbohydr Research	HCAPLUS
Kragen	11983	[1	US 4369125	HCAPLUS
Maury	1982	ļ	1	US 4352882	HCAPLUS
McNeely	1968	[}	US 3391060	HCAPLUS
Montezinos	1998	[1	US 5792502	HCAPLUS
Patton	1962	[1	US 3020206	HCAPLUS
Patton	1962	[l	US 3020207	HCAPLUS
Richmon	1983	İ	1	US 4375512	HCAPLUS
Stankowski, J	1993	241	321	Carbohydr Research	HCAPLUS
Tait, M	1990	13	133	Carbohydr Polym	HCAPLUS

- L64 ANSWER 5 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 2000:438241 HCAPLUS
- DN 133:336764
- TI Heterotypic interactions of deacetylated xanthan with a galactomannan of high galactose substitution during synergistic gelation
- AU Goycoolea, Francisco M.; Milas, Michel; Rinaudo, Marguerite
- CS Centro de Investigacion en Alimentacion y Desarrollo, Sonora, 83000, Mex.
- SO Special Publication Royal Society of Chemistry (2000), 251 (Gums and Stabilisers for the Food Industry 10), 229-240 CODEN: SROCDO; ISSN: 0260-6291
- PB Royal Society of Chemistry
- DT Journal
- LA English
- Phys. thermo-reversible gels of deacetylated xanthan AB (DX) mixed with varying concns. of galactomannan extd. from mesquite (Prosopis spp.) seed endosperm (MSG) (M/G .apprx. 1.1; MW .apprx. 2.1 .times. 106) set and melt co-operatively at .apprx.23-27.degree.C in 5 mM NaCl. The liq.-like character of the gels at 20.degree.C (tan .delta.20.degree.C) and at the gelling temp. (tan .delta.critical), attained their min. values when the concn. of MSG was .apprx.0.4-0.5 g L-1, while holding fixed that of DX at .apprx.1.0 g L-1. Mech. elasticity (G'), increased progressively as the proportion of MSG incorporated in the mixt. increased from 0 up to .apprx.0.5 g L-1 and beyond this concn., G' values showed little further change with increasing MSG concn.; this initial behavior is related to a progressive crosslinking of galactomannan and DX leading to establishment of a temporary gel network. Both values of onset temp. of gel formation (Tg) and that of DSC midpoint thermal transition (Tm), are in good agreement (Tg .apprx.23.3.degree.C and Tm .apprx.26.0.degree.C) and vary slightly in mixts. within this range of compn. However, as the amt. of unbound MSG increased for a MSG/DX wt. ratio .gtoreq. 0.6, Tg values increased progressively up to

.apprx.27.0.degree.C, while Tm also showed a break-point in the pattern of behavior for setting and melting as a function of compn. For mixts. of MSG:DX wt. ratio < .apprx.0.5, the interaction in the system has the characteristics of an heterotypic assocn. process, resulting in the creation of a coupled gel network. The optimum stoichiometric ratio seems to involve a 1:1 polymer chain pair (based on the contour length) of high galactose galactomannan and disordered DX species. Greater Tg and Tm values, beyond the stoichiometric compn. ratio may result from the increase in solvent viscosity due to unbound surplus galactomannan.

11138-66-2, Xanthan
RL: PEP (Physical, engineering or chemical process); PROC (Process)
 (deacetylated; heterotypic interactions of
 deacetylated xanthan with a galactomannan of high
 galactose substitution during synergistic gelation)

RETABLE

ΙT

Referenced Author	Year	VOL	PG	Referenced Work	Referenced
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	+=====	+====	+=====	+===========	+=======
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Bresolin, T	1998	123	263	Int J Biol Macromol	HCAPLUS .
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Chambon, F	1987	31	683	J Rheol	HCAPLUS
Chandrasekaran, R	11997			Carbohydr Polym	HCAPLUS
Cheetham, N	11986	16	257	Carbohydr Polym	HCAPLUS
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Fernandes, P		24	1269	Biopolymers	1
Foster, T	1992	ľ		Thesis University of	
Ganter, J	1995		13	Int J Biol Macromol	HCAPLUS
Gomes, C	1998		1239	Gums and Stabilisers	HCAPLUS
Goycoolea, F	1995	128	351	Carbohydr Polym	HCAPLUS
Goycoolea, F	11995		8308	Macromolecules	HCAPLUS
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L64 ANSWER 6 OF 40 HCAPLUS COPYRIGHT 2002 ACS

AN **1999:169667** HCAPLUS

DN 130:322537

TI A stray field magnetic resonance study of water diffusion in bacterial exopolysaccharides

AU Hart, T. D.; Chamberlain, A. H. L.; Lynch, J. M.; Newling, B.; McDonald, P. J.

- CS School of Biological Sciences, University of Surrey, Guildford, GU2 5XH, UK
- SO Enzyme and Microbial Technology (1999), 24(5/6), 339-347 CODEN: EMTED2; ISSN: 0141-0229
- PB Elsevier Science Inc.
- DT Journal
- LA English
- Nuclear (1H) magnetic stray field gradient methods have been used to AΒ measure the concn. dependence of the water self-diffusion coeff. (Dself) in the com. available bacterial exopolysaccharide xanthan and a chem. derived deacetylated form. The Dself coeff. of water is interpreted to directly relate to the degree of water binding in the polysaccharide gel. The removal of acetyl groups from xanthan has been shown to result in a redn. in Dself at any given polymer concn. In addn., stray field magnetic resonance profiling (1H) has been used to measure the rate at which water diffuses through a polysaccharide gel at a range of polymer concns. (Dmutual coeff. of water) in: xanthan; deacetylated xanthan and polymers produced by the soil bacteria, Enterobacter cloacae and Azotobacter chroococcum. Samples with a reduced acetyl or uronic acid content showed a lower Dmutual coeff. at a range of polymer concns. The lower Dself coeff. found for deacetylated xanthan is believed to contribute to the lower Dmutual coeff. obtained relative to the native mol. The obsd. link between the mobility (Dself) and transport (Dmutual) of water in bacterial exopolysaccharides furthers our understanding of the role(s) of these materials for bacteria and opens new opportunities for engineering bacteria for improved survival in water-stressed environments.
- IT 11138-66-2, Xanthan gum 11138-66-2D,

Xanthan gum, deacetylated

RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process); USES (Uses)

(a stray field magnetic resonance study of water diffusion in bacterial exopolysaccharides)

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     1999:77644 HCAPLUS
ΑN
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    130:141516
ΤI
    Deacetylated xanthan gum-containing
     guar-free aqueous drilling fluids for petroleum wells
IN
    Langlois, Bruno
PA
    Rhodia Chimie, Fr.
SO
     PCT Int. Appl., 33 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     French
FAN.CNT 1
                      KIND DATE
     PATENT NO.
                                           APPLICATION NO.
                                                             DATE
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     WO 9903948
                            19990128
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                                          WO 1998-FR1514 19980710
                      A1
         W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
             DK, EE, ES, FI, GB, GE, GH, GM, HU, ID, IL, IS, JP, KE, KG, KP,
             KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO,
             NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA,
             UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI,
             CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
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             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
    NO 2000000208
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                            20000317
                                           NO 2000-208
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     US 2002137635
                       A1
                                           US 2001-82555
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                            20020926
PRAI FR 1997-9087
                       Α
                            19970717
    WO 1998-FR1514
                       W
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US 2000-462995

A1

20000114

- AB Guar-free aq. fluids for use in oil exploration contains deacetylated xanthan gum, in the form of a pentamer, which is combined with at least one compd. that increases the ionic strength of the medium and a std. filtrate reducing compd. Compds. that increase the ionic strength include org. or mineral acids, salts (e.g., halides, sulfates, carbonates, bicarbonates, silicates, phosphates, and formates), and alkali metal and alk. earth metal formates and acetates. The fluids typically contain 0.01-2 wt.% deacetylated guar gum, 5000-110,000 ppm of compds. that increase the ionic strength of the medium, and 0-1 wt.% of a filtrate-reducing Typical filtrate-reducing compds. include cellulose derivs., polyacrylamides, polyacrylates, succinoglycans, natural starches and starch derivs., and carbons. Other components present can include dispersants (e.g., polyphosphates, tannins, polynaphthalenesulfonates, etc.), oxygen scavengers, and densifying agents (e.g., zinc salts, iron oxides, barite, etc.).
- IT 11138-66-2D, Xanthan gum, deacetylated

RL: TEM (Technical or engineered material use); USES (Uses) (drilling fluids contq.; deacetylated xanthan

gum-contg. guar-free aq. drilling fluids for petroleum wells)

RETABLE

Referenced Auth (RAU)	(RPY) (RV	L) (RPG	Referenced Work G) (RWK)	Referenced File
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Vanderslice, R	1989		US 4868293 A	HCAPLUS
Wellington, S	1980		US 4218327 A	HCAPLUS

- L64 ANSWER 8 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1996:309789 HCAPLUS
- DN 125:8943
- TI Influence of acyl substituents on the interaction of xanthans with plant polysaccharides
- AU Ross-Murphy, S. B.; Shatwell, K. P.; Sutherland, I. W.; Dea, I. C. M.
- CS Division Life Sciences, King's College London, London, W8 7AH, UK
- SO Food Hydrocolloids (1996), 10(1), 117-122 CODEN: FOHYES; ISSN: 0268-005X
- PB Oxford University Press
- DT Journal
- LA English
- AB Small deformation rheol. and measurement of crit. gelling concns. has been carried out to study the interactions between solns. of microbially cultured variant xanthans and chem. modified samples of this polymer, with three plant polysaccharides, guar gum, locust bean gum and konjac mannan. Using these methods we have been able to assess the influence of the acyl substituents upon the interaction behavior.
- IT 11138-66-2, Xanthan

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process)

(acyl substituents effect on interaction of xanthans with plant polysaccharides)

- L64 ANSWER 9 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1996:309785 HCAPLUS
- DN 125:8939
- TI Release of disordered xanthan oligomers upon partial acid hydrolysis of double-stranded xanthan
- AU Stokke, Bjorn Torger; Christensen, Bjorn Erik
- CS Department Physics, University Trondheim, Trondheim, N-7034, Norway

- SO Food Hydrocolloids (1996), 10(1), 83-89 CODEN: FOHYES; ISSN: 0268-005X
- PB Oxford University Press
- DT Journal
- LA English
- Double stranded xanthan was subjected to acid hydrolysis and was AΒ found to yield xanthans with partly truncated sidechains and decreasing mol. wt. The depolymn. kinetics of xanthan deviated from that of dispersed single-stranded polymers by revealing an initial regime in which the xanthan prepn. showed only minor decrease in mol. wt. (referring to the intact polypentamer repeating unit). Subsequently, a more rapid degrdn. occurred. In the latter phase an oligomeric fraction yielding an overall bimodal mol. wt. distribution, also started to appear in the mol. wt. distribution. This oligomeric fraction had the same chem. compn. as the rest of the sample. Exptl. detn. of the mass per unit length suggests that the partly hydrolyzed xanthans also possessed the basic duplex structure of intact xanthan. These data can be accounted for by taking the cooperative nature of the dimerization occurring on pairing two xanthan chains into account in a Monte Carlo model for depolymn. of duplex polymers.
- IT 11138-66-2, Xanthan
 - RL: RCT (Reactant); RACT (Reactant or reagent)
 (double-stranded; release of disordered xanthan oligomers on
 partial acid hydrolysis of)
- L64 ANSWER 10 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1996:306131 HCAPLUS
- DN 124:346429
- TI Screening for synergistic interactions in dilute polysaccharide solutions
- AU Goycoolea, F. M.; Morris, E. R.; Gidley, M. J.
- CS Silsoe College, Silsoe, Cranfield University, Bedford, MK45 4DT, UK
- SO Carbohydrate Polymers (1996), Volume Date 1995, 28(4), 351-358 CODEN: CAPOD8; ISSN: 0144-8617
- PB Elsevier
- DT Journal
- LA English
- AB A simple viscometric approach was used to screen for binding interactions between different polysaccharides in very dil. soln. where exclusion effects should be negligible. The method involves prepg. stock solns. to approx. the same, low viscosity (.eta.sp .apprxeq. 1), dialyzing to identical ionic conditions, mixing in various proportions, and looking for departures from the initial common viscosity. Mixts. of xanthan qum or de-acetylated xanthan

qum with carob qum (I) or konjac mannan (II) show massive enhancement of viscosity, as anticipated from the formation of synergistic gels at higher concns. However, no viscosity changes upon mixing with I or II were obsd. for other conformationally ordered bacterial polysaccharides (welan and rhamsan) or for alginate and pectin with sufficient Ca2+ to induce almost complete conversion to the dimeric "egg box" form, demonstrating that conformational rigidity is not, in itself, sufficient for other polysaccharides to form heterotypic junctions with mannan or glucomannan chains. Interactions of carrageenans with I depended on both conformation and the extent of aggregation. Mixts. of I with K+ .kappa.-carrageenan (III) in 100 mM KCl (which is known to promote extensive aggregation of double helixes) gave erratic values for rotational viscosity and showed typical gel-like mech. spectra under low-amplitude oscillation. Disordered carrageenans, i.e., K+ III in water and K+ .lambda.-carrageenan in 100 mM KCl, showed no evidence of interaction with I. Neg. results were also obtained for .iota.-carrageenan (IV) under ionic conditions believed to promote ordering without significant aggregation (100 mM KCl). However, under conditions where limited aggregation might be expected, e.g. IV in 90 mM

CaCl2 and Me4N+ III in 150 mM Me4NI, significant decreases in viscosity were obsd. upon mixing with I, which may indicate some intermol. assocn. but without the formation of an extended network structure.

- L64 ANSWER 11 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- 1996:72500 HCAPLUS AN
- DN 124:261539
- Dynamic simulations of the molecular conformations of wild type and mutant TΤ xanthan polymers suggest that conformational differences may contribute to observed differences in viscosity
- ΑU Levy, Samuel; Schuyler, Scott C.; Maglothin, Ronald K.; Staehelin, L. Andrew
- Dep. Mol. Cell. Dev. Biol., Univ. Colorado, Boulder, CO, 80309-0347, USA CS
- SO Biopolymers (1996), 38(2), 251-72 CODEN: BIPMAA; ISSN: 0006-3525
- PB Wiley
- DT Journal
- LA
- English Xanthan qum is an exopolysaccharide secreted by the AΒ bacterium Xanthomonas campestris whose ability to make solns. viscous at low concns. and over a pH and temp. range have generated much interest in both academic and industrial environments. Mutant Xanthomonas strains have been derived that produce xanthan gums with an altered or variant subunit chem. structure and different measured viscosities when compared with the wild type (wt) form of the polymer. Two variant gums were targeted as potentially interesting in this study, these being the nonacetylated tetramer (natet) and the acetylated tetramer (atet), which both lack a side-chain terminal mannose residue and in one case (natet) lacks an acetate group on an internal mannose residue. Solns. of these tetrameric gums possess viscosities higher (natet) and lower (atet) than the wt gum, and therefore we have attempted to det. whether these mols. possess unique conformational preferences when compared with the wt and with each other. In this manner we can initiate an understanding of how a polysaccharide's conformation contributes to its soln. properties. The GEGOP software permits a sampling of the static and dynamic equil. states of carbohydrate mols., and this software was employed to calc. equil. states of representative oligosaccharides with chem. structures representative of xanthan-like mols. Energy minimization techniques revealed similar local min. for all three mols. Some of these min. are comprised of elongate backbone conformations (A type) in which side chains fold onto backbone surfaces. Other min. with A backbones possessed side chains in less intimate backbone contact esp. when calcns. were performed with a low dielec. const. This phenomenon was particularly pronounced in the wt mol. where an increased no. of neg. charged side-chain residues experience charge repulsion resulting in reduced side-chain-backbone contact. Metropolis Monte Carlo (MMC) dynamic simulations performed with an elevated temp. factor (1000 K) allowed a better qual. representation of conformational space than 300 K simulations. Employing a nonhierarchical cluster anal. method (population d. profile: PDP) coupled with a classification scheme, it was possible to partition resulting MMC data sets into conformational families. This anal, revealed that in simulations performed with different dielec. const. values (10, 25, and .infin.) all mols. possessed primarily A-type backbones. Less elongate, more open helical backbone forms (B, C, D, J, and Flat-a) did occur during the simulations but were populated to a lesser extent. In the natet mol. significantly open helical backbones existed (E, F, G, H, and I) that did not occur in the lower viscosity wt and atet mols. PDP clustering methods and subsequent conformational classification applied to the first residue (mannose) of the side chain permitted a detn. of side-chain orientation. Comparison of all three mols. indicated a larger population of side-chain conformational families in less direct backbone contact for the wt mol. than either of the variant mols. (natet/atet) suggesting that the side

chains in the wt are more flexible. Thus, a major conformational difference between the high viscosity natet and the lower viscosities of the wt/atet is the increased amt. of open helical backbone in the natet. In addn., the significant difference between the higher viscosity wt and the lower viscosity atet is the increase side-chain flexibility in the wt. We hypothesize that conformational differences of this kind could form a partial explanation of the obsd. differences in viscosity between these xanthan-like polymers.

IT 11138-66-2, Xanthan

RL: PRP (Properties)

(dynamics simulations of the mol. conformations of wild type and mutant **xanthans** suggest that conformational differences may contribute to obsd. differences in viscosity)

- L64 ANSWER 12 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1996:68353 HCAPLUS
- DN 124:179354
- TI Synergistic gelation of galactomannans or konjac glucomannan: Binding or exclusion?
- AU Goycoolea, F. M.; Foster, T. J.; Richardson, R. K.; Morris, E. R.; Gidley, M. J.
- CS Cranfield Institute Technology, Silsoe College, Bedford, MK45 4DT, UK
- SO Gums and Stabilisers for the Food Industry 7, [Proceedings of the International Conference], 7th, Wrexham, UK, July 1993 (1994), Meeting Date 1993, 333-44. Editor(s): Phillips, Glyn O.; Williams, Peter A.; Wedlock, David J. Publisher: IRL Press, Oxford, UK. CODEN: 62HUAF
- DT Conference
- LA English
- AB Synergistic gels of xanthan or deacetylated . xanthan with locust bean gum (LBG) or konjak mannan (KM) melt and set at .apprx.60.degree., with no thermal hysteresis. Gel-like rheol. persists to very low concns. (0.02% w/v). Gelation occurs with the xanthan component in either its ordered or disordered form, and is accompanied, with KM as co-synergist, by large enthalpy changes (.DELTA.H) Formation and melting of synergistic gels of .kappa.-carrageenan with LBG or KM, in contrast, invariably occur a few degrees above the corresponding processes for carrageenan alone. Mutual exclusion, driving formation of an interpenetrating network of LBG or KM within the carrageenan gel, would be consistent with SEM evidence, but not with double-peaking in DSC. The onset of synergistic gelation occurs when there is, on av., about one full turn of helix structure in each carrageenan chain. We therefore suggest that the most likely interpretation is that unsubstituted regions of the LBG or KM backbone bind to the helixes as they form, thus raising the temp. of ordering and gelation.
- IT 11138-66-2, Xanthan gum 11138-66-2D,

Xanthan gum, deacetylated

RL: PEP (Physical, engineering or chemical process); PROC (Process) (synergistic gelation of galactomannans or glucomannans)

- L64 ANSWER 13 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1996:68348 HCAPLUS
- DN 124:179351
- TI Xanthan polytetramer: Conformational stability as a barrier to synergistic interaction
- AU Foster, T. J.; Morris, E. R.
- CS Cranfield Institute Technology, Silsoe College, Bedford, MK45 4DT, UK
- Gums and Stabilisers for the Food Industry 7, [Proceedings of the International Conference], 7th, Wrexham, UK, July 1993 (1994), Meeting Date 1993, 281-9. Editor(s): Phillips, Glyn O.; Williams, Peter A.; Wedlock, David J. Publisher: IRL Press, Oxford, UK. CODEN: 62HUAF

- DT Conference
- LA English
- AB The genetically engineered polytetramer variant of xanthan, in which the terminal mannose residues of each side chain are absent, shows a thermoreversible conformational transition in soln. (as monitored by DSC, · CD, and optical rotation). As found for xanthan, there is no detectable thermal hysteresis on cooling, and 1/Tm decreases linearly with In I (where Tm is the transition midpoint temp. and I is ionic strength), but the Tm values for polytetramer are substantially higher. Native polytetramer shows normal soln. rheol., but develops some xanthan -like "weak-gel" properties after heating and cooling through the temp. range of the conformational transition. We suggest that the polymer, as biosynthesized, is fully ordered along each chain, but renatures with shorter ordered sequences that promote network formation. The polytetramer shows no evidence of synergistic assocn. with locust bean gum or konjac glucomannan, in contrast to xanthan and (particularly) deacetylated xanthan, which gave massive viscous interactions even in very dil. soln. The difference is attributed to the enthalpic stability of the polytetramer helix (.DELTA.H .sum. 8.5 J/g, in comparison with .apprx.4.0 J/g for xanthan) preventing conformational rearrangement into heterotypic junctions. Nonacetylated polytetramer (also genetically engineered) gave a similar high value of .DELTA.H and again showed no evidence of synergistic interaction. As found for xanthan, the absence of acetate groups lowers Tm, indicating that they contribute to the stability of the ordered structure.
- IT 11138-66-2D, Xanthan gum, polytetramer derivs.

RL: PRP (Properties)

(conformational stability as a barrier to synergistic interaction)

- L64 ANSWER 14 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1995:894421 HCAPLUS
- DN 123:312460
- TI Stoichiometry and Conformation of Xanthan in Synergistic Gelation with Locust Bean Gum or Konjac Glucomannan: Evidence for Heterotypic Binding
- AU Goycoolea, F. M.; Richardson, R. K.; Morris, E. R.; Gidley, M. J.
- CS Silsoe College, Cranfield University, Silsoe/ Bedford, BEDFORD, UK
- SO Macromolecules (1995), 28(24), 8308-20 CODEN: MAMOBX; ISSN: 0024-9297
- PB American Chemical Society
- DT Journal
- LA English

AB

Synergistic gels of xanthan or deacetylated xanthan (DX) with locust bean gum (LBG) or konjac glucomannan (KM) melt and set at .apprx.60 .degree.C, with no thermal hysteresis. Gelation occurs with the xanthan component in either its ordered or its disordered form and, with KM as cosynergist, is accompanied by large enthalpy changes (.DELTA.H) in DSC. Gel modulus (G') and .DELTA.H increase linearly with increasing ratio of KM:DX up to .apprx.1:1, with little further change at higher ratios. Liq.-like character (tan .delta.) passes through a sharp min. at about the same Mixed gels of KM with unmodified xanthan show similar behavior, but the max. value of .DELTA.H is lower, and the proportion of KM required to achieve this max. is higher. The heat changes (per gram of xanthan or DX) depend only on mixing ratio, not on total concn., arguing strongly for stoichiometric binding rather than an exclusion mechanism. With LBG in place of KM, the sol-gel transition is much wider and gives no discernible peaks in DSC. The min. in tan .delta. with varying compn., however, is still evident, again arguing for a binding process, and the moduli are higher (.apprx.3.times.). Gels incorporating KM show evidence of structural rearrangement after their initial formation (maxima in the temp. dependence of G''; shoulders in tan .delta. and in

DSC); no such effects are seen for LBG. In the light of previous X-ray diffraction studies in the condensed phase, it is suggested that initial gelation involves heterotypic junctions between **xanthan** or DX and KM or LBG, with both components in a 21 conformation, but that junctions involving KM convert to a more compact 6-fold arrangement at lower temp.

IT 11138-66-2, Xanthan 11138-66-2D,

Xanthan, deacetylated

RL: BSU (Biological study, unclassified); PEP (Physical, engineering or chemical process); BIOL (Biological study); PROC (Process) (stoichiometry and conformation of xanthan in synergistic gelation with locust bean gum or konjac glucomannan)

- L64 ANSWER 15 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1995:708226 HCAPLUS
- DN 123:314302
- TI Synergistic interaction between **xanthan** and galactomannan isolated from Leucaena leucocephala de Wit
- AU Pakdee, Parwadee; Tako, Masakuni; Yokohari, Tetsuo; Kinjyo, Kazuhiko; Hongo, Hujiya; Yaga, Shiryo
- CS Coll. Agric., Univ. Ryukyus, Okinawa, 903-01, Japan
- SO Oyo Toshitsu Kagaku (1995), 42(2), 105-13 CODEN: OTKAE3; ISSN: 1340-3494
- PB Nippon Oyo Toshitsu Kagakkai
- DT Journal
- LA English
- AΒ The non-Newtonian behavior and dynamics viscoelasticity of a series of ag. mixts. of xanthan and galactomannan isolated from Leucaena leucocephala de Wit were measured with a rheologiometer. At a concn. of 0.2% of total qums, gelation did not occur at room temp., but at a low temp. (0.degree.). A much stronger interaction was obsd. with mixts. contq. deacetylated, deacylated, or native xanthan than with depyruvated xanthan. The max. dynamic modulus was obtained when the ratio of xanthan to galactomannan was 2:1. The dynamics viscoelasticity parameters for mixts. with deacetylated and native xanthan decreased rapidly at temps. above 20 and 15.degree., resp. It was concluded that the side chains of the galactomannan mol. prevent intermol. interaction between xanthan and galactomannan. The results obtained support the interaction mechanism between xanthan and locust-bean gum previously proposed.
- IT 11138-66-2, Xanthan

RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent) (synergistic interaction between xanthan and galactomannan isolated from leucaena leucocephala de wit)

- L64 ANSWER 16 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1995:409865 HCAPLUS
- DN 122:182878
- TI Studies on a viscous, gel-forming exopolysaccharide from Sphingomonas paucimobilis GS1
- AU Ashtaputre, Anita A.; Shah, Avinash K.
- CS Dep. Microbiol. Biotechnol. Cent., M. S. Univ. Baroda, Baroda, 390 002, India
- SO Applied and Environmental Microbiology (1995), 61(3), 1159-62 CODEN: AEMIDF; ISSN: 0099-2240
- PB American Society for Microbiology
- DT Journal
- LA English
- AB A new strain, Sphingomonas paucimobilis GS1, accumulated 6.5 g of a highly viscous exopolysaccharide per L, using sucrose as a substrate. The anionic heteropolysaccharide contained the following, in grams per g: glucose, 0.7; galacturonic acid, 0.11; glucuronic acid, 0.07; and acetate,

0.12. The viscosity of the exopolysaccharide (4.0~g/L;~4,200~cP) was 5.5 times that of xanthan gum and was stable over a wide pH and temp. range as well as in the presence of NaCl. Deacetylated polymer produced a clear, agarlike, thermoreversible gel in the presence of cations. The gel strength of the modified polymer was four times that of agar and could withstand autoclaving.

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L64 ANSWER 17 OF 40 HCAPLUS COPYRIGHT 2002 ACS
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- AN 1992:614877 HCAPLUS
- DN 117:214877
- TI Synergistic interaction between **xanthan** and konjac glucomannan in aqueous media
- AU Tako, Masakuni
- CS Dep. Biosci. Biotechnol., Univ. Ryukyus, Nishihara, 903-01, Japan
- SO Bioscience, Biotechnology, and Biochemistry (1992), 56(8), 1188-92
- DT Journal
- LA English
- AB The non-Newtonian behavior and dynamic modulus of a series of aq. mixts. of xanthan (native, deacetylated, depyruvated, and deacylated) and konjac glucomannan were measured with a rheogoniometer. The flow curves, at 55.degree., of a mixed soln. of xanthan and glucomannan showed plastic behavior at 0.1% total gums. At a concn. of 0.1% total gums, gelation occurred at room temp. A much stronger gel was obsd. in a mixt. with deacetylated xanthan, i.e., about twice as strong as that of a mixt. with depyruvated xanthan. The dynamic modulus of a mixt. of deacylated xanthan and glucomannan stayed at very small value in the presence of CaCl2 (6.8 mM) and urea (4.0 M). The side chains of xanthan were dominant in the interaction with konjac glucomannan mols.
- IT 11138-66-2, Xanthan gum 11138-66-2D,

CODEN: BBBIEJ; ISSN: 0916-8451

Xanthan gum, deacylated

RL: USES (Uses)

(synergistic interactions with konjac glucomannan in aq. media)

- L64 ANSWER 18 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1992:61950 HCAPLUS
- DN 116:61950
- TI Role of conformation and acetylation of xanthan on xanthan-guar interaction
- AU Lopes, L.; Andrade, C. T.; Milas, M.; Rinaudo, M.
- CS Inst. Macromol., Univ. Fed. Rio de Janeiro, Rio de Janeiro, 20000, Brazil
- SO Carbohydrate Polymers (1991), Volume Date 1992, 17(2), 121-6 CODEN: CAPOD8; ISSN: 0144-8617
- DT Journal
- LA English
- AB The synergistic effect obtained by mixing xanthan and guar solns. were examd. by low-shear viscosity measurements in relation to the temp. Native and deacetylated xanthan samples were used in mixts. in which the total polymer concns. were 1 g/L and 0.5 g/L. Gelation was obsd. for temps. <15.degree. for the native xanthan -guar system (wt. ratio 1/1) in 10-2 M NaCl and at 22-24.degree. for the same system in water; in this last case, it is known that the xanthan is in the disordered conformation. For a mixt. of deacetylated xanthan-guar, gelation was obsd. below 26.degree. in water. There was a stronger interaction between deacetylated xanthan and guar than native xanthan and guar because of enhanced xanthen-guar gum backbone assocn. in the former case.
- IT 11138-66-2, Xanthan gum RL: PRP (Properties)

(acetylation and conformation of, guar gumxanthan gum interactions in relation to)

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ANSWER 19 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
AN
    1991:635120 HCAPLUS
DN
    115:235120
ΤI
     Synergistic interaction between deacylated xanthan and
     galactomannan
ΑU
     Tako, Masakuni
     Dep. Agric. Chem., Univ. Ryukyus, Nishihara, 903-01, Japan
CS
SO
     Journal of Carbohydrate Chemistry (1991), 10(4), 619-33
     CODEN: JCACDM; ISSN: 0732-8303
DT
     Journal
LA
     English
AB
    The dynamic modulus and optical rotation of a mixed soln. of denatured -
    xanthan (depyruvated and deacylated) and galactomannan
     (locust-bean gum and guar gum) were measured with a
     rheogoniometer and a polarimeter. Gelation occurred in a mixt. of native
    xanthan with locust-bean gum at a concn. of 0.2% total
     gums at room temp., but not with guar gum. A mixt. of
     deacylated xanthan and locust-bean gum showed
     the highest dynamic modulus, .apprx.3 times as strong as that of a mixt.
     with depyruvated xanthan. The dynamic modulus of a mixt. of
     deacylated xanthan and locust-bean gum stayed
     at very small value in the presence of CaCl2 (6.8 mM) and urea (4.0 M).
     Possible binding sites between deacylated xanthan and
     locust-bean qum mols. are proposed.
ΙT
     11138-66-2D, Xanthan gum, deacylated
     RL: USES (Uses)
        (galactomannan blends, gelation of, synergism in)
    ANSWER 20 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
AN
     1991:536533 HCAPLUS
DN
     115:136533
     Hydrolysis of xanthan in dilute acid: effects on chemical
ΤI
     composition, conformation, and intrinsic viscosity
ΑU
     Christensen, Bjoern E.; Smidsrod, Olav
CS
     Norweg. Biopolym. Lab., Univ. Trondheim, Trondheim, N-7034, Norway
SO
     Carbohydrate Research (1991), 214(1), 55-69
     CODEN: CRBRAT; ISSN: 0008-6215
DT
     Journal
     English
LA
    The polysaccharide xanthan has been depolymd. by mild acid
AB
     hydrolysis (pH 1-4) at 80.degree.. The conformational state was varied
     from fully ordered to partially disordered by varying the ionic strength
             Hydrolysis occurred mainly in the side chains, with the terminal
     .beta.-mannose as the most susceptible unit, yielding a continuous series
     of modified xanthans from the intact "polypentamer" to
     the "polytetramer", while retaining a high mol. wt. Depolymn. of the
     glucan backbone was analyzed by monitoring the intrinsic viscosity.
     ordered xanthan the calcd. changes in the degree of polymn. as a
     function of time deviate strongly from that expected for random depolymn.
     of a single-stranded, linear polymer, but the data are in qual. agreement
     with the behavior of such double-stranded polymers as DNA.
     formational properties of partly hydrolyzed xanthan were
     investigated by optical rotation.
IT
     11138-66-2, Xanthan
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydrolysis of, in dil. acid, conformation and intrinsic viscosity
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L64 ANSWER 21 OF 40 HCAPLUS COPYRIGHT 2002 ACS AN 1991:516704 HCAPLUS

effect on)

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DN
    115:116704
ΤI
     Synergistic interaction between xanthan and tara-bean
ΑU
     Tako, Masakuni
     Dep. Agric. Chem., Univ. Ryukyus, Nishihara, 903-01, Japan
CS
SO
    Carbohydrate Polymers (1991), 16(3), 239-52
    CODEN: CAPOD8; ISSN: 0144-8617
DT
     Journal
LA
     English
    The non-Newtonian behavior and dynamic viscoelasticity of a series of aq.
AΒ
    mixts. of xanthan and tara-bean gums were measured
     with a rheogoniometer. At a concn. of 0.2% of the total gum,
     gelation did not occur at room temp., but rather at a low temp.
     (0.degree.). A much stronger interaction was obsd. with mixts. contg.
     deacetylated, deacylated, or native xanthan
     than with depyruvated xanthan. The max. dynamic modulus was
     obtained at a xanthan-tara-bean gum ratio of 1:2. The
     dynamic viscoelastic parameters for mixts. with deacetylated and
     deacylated xanthan decreased rapidly at temps. above 25
     and 20.degree., resp. The side chains of the tara-bean gum mol.
    prevented an intermol. interaction between xanthan and tara-bean
    gums.
IT
     11138-66-2, Xanthan gum
     RL: USES (Uses)
        (tara-bean gum mixts., viscoelasticity of, synergism in)
     11138-66-2D, Xanthan gum, deacylated
TT
     RL: USES (Uses)
        (tara-bean qum mixts., viscoelasticity of, synergism in
       relation to)
    ANSWER 22 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
ΑN
    1991:45353 HCAPLUS
DN
    114:45353
TI
     Influence of the acetyl substituent on the interaction of
    xanthan with plant polysaccharides. III. Xanthan
     -konjac mannan systems
ΑU
     Shatwell, Karolyn P.; Sutherland, Ian W.; Ross-Murphy, Simon B.; Dea, Iain
    C. M.
CS
     Dep. Microbiol., Sch. Agric., Edinburgh, EH9 3JG, UK
SO
    Carbohydrate Polymers (1990), 14(2), 131-47
    CODEN: CAPOD8; ISSN: 0144-8617
DT
     Journal
LA
     English |
AB
    A range of xanthans (Na+ salt form) with varying levels of
    acetyl and pyruvic acid substitution were prepd. by culturing
     different strains of Xanthomonas campestris and by chem.
     deacetylation and depyruvylation. Oscillatory-shear measurements
     were used to characterize the interaction between these polymers and
     konjac mannan in deionized water and the data was analyzed statistically.
     The majority of the polymers interacted to form a strong
     thermoreversible-gel network. Xanthan gelled with konjac mannan
     only at relatively high concns. compared with xanthan-locust
     bean gum (LBG) systems but the gels formed had significantly
     higher melting and setting temps. than xanthan-LBG gels of the
     same concn. The transition from the liq. to the gel state was also much
     sharper.
              The strength of the gels was heavily dependent on the level of
     acetyl substitution.
ΙT
     11138-66-2, Xanthan gum
    RL: PRP (Properties)
        (interaction of, with Konjac mannan, acetyl substitution
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effect on)

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AN
     1991:45352 HCAPLUS
DN
     114:45352
ΤI
     Influence of the acetyl substituent on the interaction of
     xanthan with plant polysaccharides. II. Xanthan-guar
     gum systems
ΑU
     Shatwell, Karolyn P.; Sutherland, Ian W.; Ross-Murphy, Simon B.; Dea, Iain
     C. M.
     Dep. Microbiol., Sch. Agric., Edinburgh, EH9 3JG, UK
CS
SO
     Carbohydrate Polymers (1990), 14(2), 115-30
     CODEN: CAPOD8; ISSN: 0144-8617
DΨ
     Journal
LA
    English
    A range of xanthans (Na+ salt form) with varying levels of
AB
     acetyl and pyruvic acid substitution were prepd. by culturing
     different strains of Xanthomonas campestris and by chem.
     deacylation and depyruvylation. Oscillatory-shear measurements
     were used to characterize the behavior of xanthan and guar
     gum alone, and of mixts. of the 2 in deionized water; the
     xanthan under these conditions was largely in the disordered form.
     The mech. spectra of the blends resembled an entanglement network system
     and showed some features characteristic of the individual components.
     However, evidence from both rheol. and chirooptical measurements indicated
     a possible weak interaction between some low-acetyl
    xanthans and guar.
ፐጥ
    11138-66-2, Xanthan gum
     RL: PRP (Properties)
        (interaction of, with guar gum, acetyl substitution
       effect on)
    ANSWER 24 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
ΑN
     1990:631852 HCAPLUS
DN
     113:231852
TΙ
     The influence of acetyl and pyruvate substituents on the
     helix-coil transition behavior of xanthan
ΑU
     Shatwell, Karolyn P.; Sutherland, Ian W.; Dea, Iain C. M.; Ross-Murphy,
     Simon B.
CS
     Dep. Microbiol., Sch. Agric., Edinburgh, EH9 3JG, UK
SO
     Carbohydr. Res. (1990), 206(1), 87-103
     CODEN: CRBRAT; ISSN: 0008-6215
     Journal
DT
LA
    English
AB
    Xanthans (Na+ salt form) having various contents of
     acetyl and pyruvic acid groups were prepd. by culturing different
     strains of Xanthomonas campestris and by deacetylation and
     depyruvylation. Optical rotation ([.alpha.]365) was used to characterize
     the helix-coil transition behavior of these polymers in deionized water.
     There were correlations between the acetyl and pyruvic acid
     contents and the mid-point temp. of the transition, between the pyruvic
     acid content and [.alpha.]365 in the high-temp.-plateau (coil) region of
     the curve, and between the content of pyruvic acid and the height of the
     transition. In deionized water, each of the polymers showed marked
     thermal hysteresis and a time-dependent fall in [.alpha.]365, at low
            This behavior, which was attributed to kinetic factors, was
     eliminated by the addn. of NaCl. Salt also increased the melting temp.
     and reduced [.alpha.]365 in the low-temp.-plateau region of the curve in
     relation to the charge carried by the polymer. A high-pyruvate, low-
     acetyl xanthan exhibited unusual two-phase helix-coil
     transition behavior in the presence of salt.
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IT 11138-66-2, Xanthan gum RL: PRP (Properties)

(helix-coil transition of, acetyl and pyruvate group effect

- L64 ANSWER 25 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1990:424384 HCAPLUS
- DN 113:24384
- TI Influence of acetyl and pyruvate substituents on the solution properties of xanthan polysaccharide
- ΑU Shatwell, Karolyn P.; Sutherland, Ian W.; Ross-Murphy, Simon B.
- Dep. Microbiol., Univ. Edinburgh, Edinburgh, EH9 3JG, UK CS
- SO Int. J. Biol. Macromol. (1990), 12(2), 71-8 CODEN: IJBMDR; ISSN: 0141-8130
- DTJournal
- LA English
- AΒ Xanthan, an exocellular polysaccharide produced by the plant pathogenic bacterium Xanthomonas campestris has been the subject of considerable interest in recent years because of its unusual rheol. properties in soln. (weak gel) and consequent range of application. polymer consists of a cellulosic backbone with trisaccharide side chains linked to alternate backbone residues; acetyl and pyruvate substituents are carried in variable amts. on these side chains. of xanthans differing in the percentage of substituent groups and in mol. wt. range were prepd. by culturing a variety of different strains of X. campestris. All of the xanthans were characterized by a range of physicochem. techniques. In particular, the intrinsic viscosities at low shear rates, and at a range of ionic strengths, were detd. and the geometric persistence lengths evaluated by the Smidsrod-Haug method. Intensity light scattering measurements were made using the procedure of Coviello and co-workers to promote mol. dispersion. Despite significant differences in the acetyl and pyruvate contents, the mol. wt. vs mean square radius behavior of the samples did not differ substantially from each other or from those reported for other xanthan samples in the literature. The persistence length, detd. by the method of Schmidt et al. (120 .+-. 8 nm) was also, within exptl. error, the same for all the samples measured. These values differed considerably from those calcd. from the ionic strength dependence of intrinsic viscosity (the Smidsrod-Haug method) as reported by Tinland and Rinaudo and calcd. for these samples. illustrates the limitations of the latter method when applied to systems where the electrostatic contribution to the persistence length is only a small fraction of the geometrical contribution. The values obtained from the light scattering measurements support other recent conclusions that the inherent stiffness of the xanthan macromol. is not greatly influenced by the pattern of acyl substitution.
- 11138-66-2, Xanthan IT
 - RL: PRP (Properties)
 - (soln. properties of, acetyl and pyruvate substituents effect
- ANSWER 26 OF 40 HCAPLUS COPYRIGHT 2002 ACS L64
- ΑN 1989:595280 HCAPLUS
- DN 111:195280
- ΤI Evidence for intramolecular associations in xanthan molecules in aqueous media
- Tako, Masakuni; Nakamura, Sanehisa IIA
- Coll. Agric., Univ. Ryukyus, Nishihara, 903-01, Japan CS
- SO Agric. Biol. Chem. (1989), 53(7), 1941-6 CODEN: ABCHA6; ISSN: 0002-1369
- DΤ Journal
- LA English
- AB The non-Newtonian behavior and dynamic viscoelasticity of deacylated xanthan soln. were measured with a rheogoniometer. The flow curves for the deacylated xanthan at concns. below 0.5% approximated to shear-thinning behavior, and a plastic behavior above 0.8%. The apparent viscosity of a 0.5% soln. of deacylated xanthan increased with

increasing temp. up to 25, which was estd. to be a transition temp., then it decreased rapidly. The dynamic modulus of deacylated xanthan increased a little with increase of temp. up to 25 and 30.degree. in 0.8 and 1.0% solns., then it decreased rapidly with further increases of temp. The sp. rotation of deacylated xanthan stayed const. up to 25.degree., then it decreased rapidly with further increases in temp. Possible mode of intramol. assocn. between an alternate hydroxyl group at C(3) and the adjacent hemiacetal O atom of the D-glucosyl residues, and between the Me group of the acetyl residue and the adjacent hemiacetal O atom of the D-glucosyl residue were proposed.

IT 11138-66-2, Xanthan RL: PRP (Properties)

(intramol. assocn. in, in aq. media)

- L64 ANSWER 27 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1988:473771 HCAPLUS
- DN 109:73771
- TI Rheological properties of depyruvated xanthan in aqueous media
- AU Tako, Masakuni; Nakamura, Sanehisa
- CS Coll. Agric., Univ. Ryukyus, Nishihara, 903-01, Japan
- SO Agric. Biol. Chem. (1988), 52(6), 1585-6 CODEN: ABCHA6; ISSN: 0002-1369
- DT Journal
- LA English
- AB The viscosity and viscoelasticity of depyruvated xanthan in water were characterized. Depyruvated xanthan was found to have fewer intermol. assocns. than native or deacetylated xanthans that contained pyruvate.
- IT 11138-66-2, Xanthan gum

RL: PRP (Properties)

(viscoelasticity and viscosity of, in aq. soln., role of pyruvate interactions in)

IT 11138-66-2D, Xanthan gum, depyruvated deriv.

RL: PRP (Properties)

(viscosity and viscoelasticity of, in aq. soln.)

- L64 ANSWER 28 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1988:468188 HCAPLUS
- DN 109:68188
- TI Cloning and sequencing of xanthomonas campestris DNA encoding xanthan gum biosynthetic enzymes for use in manufacture of xanthan gums by recombinant bacteria
- IN Capage, Michael A.; Doherty, Daniel H.; Betlach, Michael R.; Vanderslice, Rebecca W.
- PA Getty Scientific Development Co., USA
- SO PCT Int. Appl., 152 pp.

CODEN: PIXXD2

- DT Patent
- LA English
- FAN.CNT 1

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    US 1987-29530
                           19870323
    US 1986-844332
                           19860326
    WO 1987-US604
                           19870324
    US 1988-188687
                           19880427
                           19890403
    US 1989-333868
    US 1992-815615
                           19920107
AB
    A gene cluster encoding enzymes necessary for the biosynthesis of
    xanthan gum is isolated from X. campestris and the DNA
    is sequenced. Plasmids contg. these genes can be transferred to bacteria
    which can be grown anaerobically and/or at temps. >30.degree. thus
    allowing a more economical manuf. of xanthan gum. The
    genes and the enzymic activity of the corresponding proteins were
    identified by Tn10 mutagenesis and subsequent anal. of lipid-linked
    radioactive precursors of xanthan gum. Plasmid
    pRK290-H336, which contains the gum gene cluster and has a broad host
    range, was conjugally transferred to Pseudomonas stutzeri (from
    Escherichia coli). The expression of one of the enzymes, Transferase III,
    was detd. by Western immunoblots to be almost equiv. to that in X.
    campestris itself.
TT
    11138-66-2DP, Xanthan gum, non-pyruvylated
    and/or non-acetylated 11138-66-2P,
    Xanthan gum
    RL: PREP (Preparation)
        (manuf. of, with recombinant bacteria, improved culture conditions in
       relation to)
    ANSWER 29 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
    1988:185289 HCAPLUS
AN
    108:185289
DN
TI
    Manufacture of xanthan gums with altered
    acetylation and/or pyruvylation using Xanthomonas compestris
    mutants or lysates of these mutants
    Doherty, Daniel H.; Ferber, Donna M.; Marrelli, John D.; Vanderslice,
IN
    Rebecca W.
PA
    Getty Scientific Development Co., USA
SO
    PCT Int. Appl., 37 pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 4
    PATENT NO.
                     KIND DATE
                                        APPLICATION NO. DATE
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                                          -----
    WO 8705939
                    A1
                           19871008
PΙ
                                          WO 1987-US606 . 19870324 <--
        W: DK, FI, JP, NO
        RW: AT, BE, CH, DE, FR, GB, IT, LU, NL, SE
    JP 63503198
                     Т2
                           19881124
                                          JP 1987-502340
                                                          19870324 <--
    JP 2559437
                      B2
                           19961204
                      A1
                           19890719
                                          EP 1987-902919
    EP 323952
                                                          19870324 <--
    EP 323952
                     В1
                           19930811
        R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE
                     A2
                                          EP 1992-111024
                                                           19870324 <--
    EP 511690
                           19921104
    EP 511690
                      AЗ
                           19921119
                           19970730
    EP 511690
                     В1
        R: AT, BE, DE, FR, GB, IT, LU, NL, SE
                                      AT 1987-902919
                                                           19870324 <--
    AT 92965
                    Ε
                          19930815
                      A2
                                          EP 1996-119095
    EP 765939
                           19970402
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    EP 765939
                     A3
                           19970514
        R: AT, BE, DE, FR, GB, IT, LU, NL, SE
                    A2 19970708
                                     JP 1996-70588
    JP 09176205
                                                          19870324 <--
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CA 1339113

A1

19970729

CA 1987-532838

19870324 <--

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AT 156190
                       Ε
                            19970815
                                           AT 1992-111024
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                       Α
    NO 8704846
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                                           NO 1987-4846
                                                             19871120 <--
                       В
    NO 172945
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    NO 172945
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                       Α
    NO 9201787
                            19880125
                                           NO 1992-1787
                                                             19920506 <--
                       Α
    US 5514791
                            19960507
                                           US 1994-232416
                                                             19940425 <--
                       Α
    US 5948651
                            19990907
                                           US 1995-406804
                                                             19950320 <--
                       B1
     US 6316614
                            20011113
                                           US 1995-475823
                                                             19950607 <--
                       A1
     US 2002103370
                            20020801
                                           US 2001-986803
                                                             20011113 <---
PRAI US 1986-842945
                       A
                            19860324
                                      <--
    US 1986-844435
                       Α
                            19860326
                                      <--
    US.1987-29090
                       Α
                            19870323
                                      <--
    US 1985-762878
                       A2
                            19850806
                                      <--
    EP 1987-902919
                       Α
                            19870324
                                      <--
    EP 1992-111024
                       A3
                            19870324
                                      <--
    JP 1987-502340
                       A3
                            19870324
                                      <--
    NO 1987-4846
                       Α1
                            19870324
                                      <--
    WO 1987-US606
                       W
                            19870324
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    US 1989-384621
                       B2
                            19890725
                                      <--
    US 1990-566875
                       В2
                            19900613
                                      <--
    US 1991-696732
                       В1
                            19910507
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    US 1992-928726
                       В1
                            19920813
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    US 1994-232416
                       A3
                            19940425
                                      <--
    US 1995-475823
                       Α3
                            19950607
                                      <--
AΒ
    X. campestris Mutants are prepd. which are deficient in an enzyme of the
    xanthan gum biosynthetic pathway (i.e. ketalase or
    acetylase) and are used to prep. non-acetylated
    and under- or non-pyruvylated xanthan qum. X.
    campestris X1006, contg. a Tn10-inactivated acetylase, and
    mutant X921, contg. a Tn10-inactivated ketalase, were grown overnight at
     30.degree. in broth contq. 2% glucose. The xanthan gums
    were prepd. by pptn. with org. solvents from the cell-free medium.
    confirmed that the former produced non-acetylated, and
    the latter non-pyruvylated xanthan gum. The
    non-pyruvylated gum has low viscosity at fermn. temp. but
     viscosity essentially equiv. to wild-type gum at higher temp.
     (80.degree.). The non-acetylated gum
    produces solns. of greater viscosity than either wild-type or com.
     available, chem.-deacetylated xanthan gum at
     comparable concns.
TΤ
     11138-66-2DP, non-acetylated or under- or
    non-pyruvylated
    RL: BMF (Bioindustrial manufacture); BIOL (Biological study); PREP
     (Preparation)
        (manuf. of, by Xanthomonas campestris mutants)
    ANSWER 30 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
    1988:167837 HCAPLUS
ΑN
DN
    108:167837
     Influence of acetyl and pyruvate contents on rheological
TΙ
     properties of xanthan in dilute solution
ΑU
     Callet, Francoise; Milas, Michel; Rinaudo, Marguerite
CS
    Cent. Rech. Macromol. Veg., Univ. Sci., Technol. Med. Grenoble,
     Saint-Martin d'Heres, 38402, Fr.
SO
     Int. J. Biol. Macromol. (1987), 9(5), 291-3
     CODEN: IJBMDR; ISSN: 0141-8130
DT
     Journal
LA
     English
AΒ
     The role of acetyl and pyruvate groups on rheol. properties of
     xanthan in dil. solns. was investigated. For this purpose, a
     series of xanthan derivs. were prepd. by chem. hydrolysis from
```

the same original sample, i.e. acetyl-free, pyruvate-free,

acetyl and pyruvate-free xanthans. Conformational

transitions of the 4 samples (native and modified xanthans) were followed by measuring optical rotation as a function of temp. Values of midpoint transition (Tm) thus obtained indicate that acetyl groups have a stabilizing effect on the ordered form of xanthan, whereas pyruvate groups have an opposite effect. After partial depolymn. by sonication, viscosities of the 4 samples were studied as a function of polymer concn. (below overlap concn. C*) and mol. wt. Unique curves were obtained for the abs. viscosity [.eta.] vs. mol. wt. and specific viscosity vs. the overlap parameter C[.eta.] for the 4 samples. This result shows that acetyl and pyruvate contents have no influence either on xanthan dil. soln. viscosity or on its intrinsic viscosity at a given mol. wt.

IT 11138-66-2D, deacetylated and depyruvylated RL: PRP (Properties) (rheol. behavior and conformation of)

IT 11138-66-2, Xanthan

RL: PRP (Properties)

(rheol. properties and conformation of, acetyl and pyruvate groups effect on)

- L64 ANSWER 31 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1986:572914 HCAPLUS
- DN 105:172914
- TI D-Mannose-specific interaction between **xanthan** and D-galacto-D-mannan
- AU Tako, Masakuni; Nakamura, Sanehisa
- CS Dep. Agric. Chem., Univ. Ryukyus, Okinawa, 903-01, Japan
- SO FEBS Lett. (1986), 204(1), 33-6 CODEN: FEBLAL; ISSN: 0014-5793
- DT Journal
- LA English
- AB A gelation occurred in a mixed soln. of xanthan and locust-bean gum at room temp.; in contrast, gelation did not occur in a soln. of xanthan and guar gum. The max. dynamic modulus was obtained when the mixing ratio of xanthan and locust-bean gum was 1:2 at 0.2% total gums. A mixt. of deacetylated xanthan and locust-bean gum showed the highest dynamic modulus, about twice that of the mixt. of native xanthan. The intermol. interaction between xanthan and locust-bean gum might occur between the side chains of the former and back-bone of the latter mols. in a lock-and-key arrangement.
- IT 11138-66-2

RL: PRP (Properties)

(interaction of, with polysaccharide gums)

- L64 ANSWER 32 OF 40 HCAPLUS COPYRIGHT 2002 ACS
- AN 1986:535845 HCAPLUS
- DN 105:135845
- TI Reactions in dispersions of different phases
- IN Fujishige, Norinaga; Numajiri, Rikio; Iegi, Masahiro
- PA Agency of Industrial Sciences and Technology, Japan; Kashima Oil Co., Ltd.
- SO Jpn. Kokai Tokkyo Koho, 6 pp. CODEN: JKXXAF
- DT Patent
- LA Japanese
- FAN.CNT 1

	PATENT NO.		DATE	APPLICATION NO.	DATE		
PI	JP 61120802 JP 01033481	A2 B4	19860607 19890713	JP 1984-241791	19841116 <		

AB Polymers (100 parts) forming viscous aq. solns. are dispersed or dissolved in >100 parts org. polar solvents and 20-300 parts water, dispersed in

nonpolar solvents to form water-in-oil-type emulsions, and allowed to react with reagents. The reactions are alkoxylation, crosslinking, deacetylation, acetalization or ketalization, and hydrolysis. Thus, xanthan gum reacted with epichlorohydrin to give a product having degree of substitution 2.0. 11138-66-2

TT

RL: RCT (Reactant)

(reaction of, with epichlorohydrin, in water-in-oil emulsions)

ANSWER 33 OF 40 HCAPLUS COPYRIGHT 2002 ACS L64

1985:202772 HCAPLUS AN

DN. 102:202772

TΙ Synergistic interaction between xanthan and guar gum

ΑU Tako, Masakuni; Nakamura, Sanehisa

CS Dep. Agric. Chem., Univ. Ryukyus, Okinawa, 903-01, Japan

SO Carbohydr. Res. (1985), 138(2), 207-13 CODEN: CRBRAT; ISSN: 0008-6215

DT Journal

LA English

AR The non-Newtonian behavior and dynamic viscoelasticity of a series of aq. mixts. of xanthan [11138-66-2] and guar gum At 0.2% of total [9000-30-0] were measured with a rheogoniometer. gums, gelation did not occur at room temp. but occurred at a low temp. (0.degree.). A much stronger interaction was obsd. with a mixt. of deacetylated xanthan than that with native xanthan. The max. dynamic modulus was obtained when the ratio of xanthan to guar gum was 2:1. The transition temps. of dynamic viscoelasticity for mixts. with native and deacetylated xanthan were obsd. at 25 and 30.degree., resp. Apparently, the side chains of the guar gum mol. prevent an intermol. interaction with the side chains of the xanthan mol. An intermol. interaction between xanthan and guar gum at low temp. might be promoted between the periphery of the side chains of the xanthan mol. and the backbone of the guar gum mol. and dissocn. takes place at the transition temp.

ΙT 11138-66-2

> RL: BIOL (Biological study) (gelation and rheol. of guar gum with)

ANSWER 34 OF 40 HCAPLUS COPYRIGHT 2002 ACS L64

ΑN 1985:94471 HCAPLUS

102:94471 DN

ΤI Rheological aspects of the intermolecular interaction between xanthan and locust bean gum in aqueous media

ΑU Tako, Masakuni; Asato, Atsushi; Nakamura, Sanehisa

Fac. Agric., Univ. Ryukyus, Okinawa, 903-01, Japan Agric. Biol. Chem. (1984), 48(12), 2995-3000 CS SO

CODEN: ABCHA6; ISSN: 0002-1369

DT Journal

English LA

Non-Newtonian behavior and dynamic viscoelasticity of a series of aq. mixed solns. of xanthan [11138-66-2] and locust bean [9000-40-2] were measured using a rheogoniometer, and the rheol. properties were analyzed. A gelation occurred in the mixt. at 0.2% total gums at room temp. The flow curves of the mixt. solns. showed a yield value and approximated to plastic behavior at 50.degree.. The max. dynamic modulus was obtained when the mixing ratio of xanthan to locust bean gum was 1:2, while comparable high moduli were also obtained in the mixing ratio of 1:3 or 1:4. A mixt. of deacetylated xanthan and locust bean gum showed the highest dynamic modulus, about 2 times that of the mixt. of native or Na-form xanthan. The dynamic modulus of the mixts. decreased rapidly with increasing temp. In contrast, the dynamic

viscosity was scarcely changed during increasing temp. in the mixing ratio of 2:1. The dynamic modulus was decreased by addn. of urea (4.0M), NaCl (0.1%), and MgCl2. Intermol. interaction between xanthan and locust bean gum might occur between the side chains of the former and backbone of the latter, as in a lock-and-key effect. ΙT 11138-66-2 RL: BIOL (Biological study) (rheol. of, locust bean gum effect on) L64 ANSWER 35 OF 40 HCAPLUS COPYRIGHT 2002 ACS AN 1985:77399 HCAPLUS 102:77399 DN Rheological properties of deacetylated xanthan in TΙ aqueous media ΑU Tako, Masakuni; Nakamura, Sanehisa Fac. Agric., Univ. Ryukyus, Okinawa, 903-01, Japan CS Agric. Biol. Chem. (1984), 48(12), 2987-93 SO CODEN: ABCHA6; ISSN: 0002-1369 DT Journal English LA AΒ Flow properties of aq. deacetylated xanthan gum solns. could be approximated to pseudoplastic behavior at <0.1% but to plastic behavior above 0.3%. The flow indexes in the power law for the deacetylated xanthan were somewhat different at various concns. The apparent viscosity of deacetylated xanthan decreased with increasing temp. at relatively low concns. from 0.1 to 0.5%, however, it increased with increasing temp., showed a max. value at 40.degree., and decreased gradually at 1.0%. Compared with native xanthan, deacetylated material showed higher dynamic viscoelasticity at high concns. The dynamic viscoelasticity of deacetylated xanthan decreased with increasing temp. at various concns. The dynamic viscoelasticity of deacetylated xanthan was decreased by addn. of urea (4.0M). This suggests that acetate residues, which are attached to the inner mannose residues of the side chains, contribute to the intramol. assocn., and that the side chains of xanthan become more flexible after deacetylation. ΤТ 11138-66-2D, deacetylated RL: BIOL (Biological study) (rheol. of) ANSWER 36 OF 40 HCAPLUS COPYRIGHT 2002 ACS L64 AN1984:612999 HCAPLUS DN 101:212999 TIHygroscopic modified polysaccharides Agency of Industrial Sciences and Technology, Japan; Kashima Oil Co., Ltd. PA SO Jpn. Kokai Tokkyo Koho, 4 pp. CODEN: JKXXAF DT Patent LA Japanese FAN.CNT 1 KIND DATE APPLICATION NO. DATE PATENT NO. -----____ -----19840815 JP 1983-16226 19830204 <--PΤ JP 59142201 A2 JP 63022201 B4 19880511 Polysaccharides deacetylated with an alkyl alcoholate or NH3 are AΒ hygroscopic. Thus, 0.12 g Na was treated with 100 mL MeOH to give 1.5 g Na methylate [124-41-4] which was reacted with xanthan gum (I) [11138-66-2] to give modified I with good hygroscopicity. TT 11138-66-2 RL: RCT (Reactant)

(deacetylation of, with sodium methylate or ammonia)

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IT
    11138-66-2DP, deacetylated
    RL: PREP (Preparation)
        (hygroscopic, manuf. of)
    ANSWER 37 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
ΑN
    1983:74177 HCAPLUS
DN
    98:74177
ΤI
    Modified xanthan - its preparation and viscosity
ΑU
    Bradshaw, I. J.; Nisbet, B. A.; Kerr, M. H.; Sutherland, I. W.
CS
    Dep. Microbiol., Edinburgh Univ., Edinburgh, EH9 3JG, UK
SO
    Carbohydr. Polym. (1983), 3(1), 23-38
    CODEN: CAPOD8
DT
    Journal
    English
LA
AB
    The hydrolysis of xanthan gum (I) [11138-66-2
    ] in soln. with (0.5%; 10 mL) 4-5 mM trifluoroacetic acid for 90 min at
    100.degree. resulted in optimum removal of pyruvic acid acetal and
    acetyl groups, and under these conditions no low mol. wt.
    carbohydrate-contq. material was released. A comparison of the viscosity
    of native and modified I in H2O and 1% KCl showed that depyruvylation and
    deacetylation have little effect on soln. viscosity of shear rates
    between 8.8 and 88.3 s-1.
IΤ
    11138-66-2
    RL: USES (Uses)
        (deacetylation and depyruvylation of, with trifluoracetic
    11138-66-2D, deacetylated, depyruvylated
TT
    RL: PRP (Properties)
        (soln. viscosity of)
    ANSWER 38 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
ΑN
    1980:551994 HCAPLUS
DN
    93:151994
TI
    Deacetylated borate-biosynthetic gum compositions
IN
    Cottrell, Ian W.; Racciato, Joseph S.
PA
    Merck and Co., Inc., USA
    U.S., 4 pp. Cont.-in-part of U.S. Ser. No. 891,575, abandoned.
SO
    CODEN: USXXAM
DT
    Patent
LA
    English
FAN.CNT 1
    PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
                                                          -----
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                                          -----
PΙ
    US 4214912
                     Α
                           19800729
                                          US 1979-45493 19790604 <--
PRAI US 1978-871279
                           19780123 <--
    US 1978-891575
                           19780330 <--
    Treatment of xanthan qum (I) with borax (II) in the
    presence of alkalies gives product with enhanced dispersibility in H2O.
    Thus, a mixt. of 6 g II, 100 mL H2O, and .apprx.19 L beer contg.
     .apprx.2.5% I was added to 150 mL 5 N NaOH and stirred for 4 h at
    40.degree. to give deacetylated I, a 1% soln. of which showed a
    viscosity of 1110 cP at pH 8.0 after storage of 2 h.
ΙT
    11138-66-2D, deacetylated
    RL: USES (Uses)
        (borax-treated, with high dispersibility in water)
    ANSWER 39 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
ΑN
    1979:591696 HCAPLUS
DN
    91:191696
TI
    Gelled compositions based on galactomannans and xanthan
PΑ
    CECA S. A., Fr.
SO
    Fr. Demande, 25 pp.
    CODEN: FRXXBL
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DT
     Patent
LA
     French
FAN.CNT 1
                     KIND DATE
                                          APPLICATION NO.
                                                            DATE
     PATENT NO.
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                           19790406
     FR 2402678
                      A1
                                          FR 1977-27467
                                                            19770912 <--
PΙ
                      В1
                           19800321
     FR 2402678
     EP 1192
                                                            19780830 <--
                      A1
                                          EP 1978-400084
                           19790321
                     В1
     EP 1192
                            19811223
        R: CH, DE, GB, NL
     US 4369125
                     Α
                            19830118
                                          US 1980-161322
                                                            19800620 <--
PRAI FR 1977-27467
                                    <--
                            19770912
     US 1978-935944
                            19780825 <--
AΒ
    Mixts. of deacetylated xanthan gum (I)
     11138-66-2] (totally or partially deacetylated) with 1
     or more galactomannan (II) [11078-30-1] at I/II ratios of 15:85 to 9:1
     formed stronger edible gels and thickening agents than did similar mixts.
     contg. regular (acetylated) xanthan gum.
     For example a I-carob qum [9000-40-2] mixt. at 30:70 gave a gel
     with rigidity .apprx.75% greater and cohesion .apprx.70% greater than of a
     gel prepd. with regular (acetylated) xanthan
     gum. The gel strength of gels contg. I was affected less by salts
     than were those contg. regular xanthan gum.
ΙT
     11138-66-2D, deacetylated
     RL: BIOL (Biological study)
        (gels contg. galactomannan and)
    ANSWER 40 OF 40 HCAPLUS COPYRIGHT 2002 ACS
L64
ΑN
     1979:543066 HCAPLUS
DN
     91:143066
    Xanthomonas biopolymer for use in the displacement of oil from
TI
     partially-exhausted deposits
IN
     Wernau, William Charles
PA
     Pfizer Inc., USA
SO
     Ger. Offen., 21 pp.
     CODEN: GWXXBX
DT
     Patent
LA
     German
FAN.CNT 1
     PATENT NO.
                      KIND DATE
                                          APPLICATION NO.
                                                            DATE
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                                          -----
                                                            -----
PI
                           19790517
                                          DE 1978-2848894
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     DE 2848894
                      Α1
     DE 2848894
                      C2
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                      Α
                                          US 1977-851757
    US 4296203
                           19811020
                                                            19771115
    DK 7803891
                      Α
                           19790516
                                                            19780904
                                          DK 1978-3891
    CA 1113875
                      A1
                          19811208
                                          CA 1978-314034
                                                            19781024
                           19790514
     BE 871955
                      A1
                                          BE 1978-191678
                                                            19781113
     GB 2008600
                      Α
                           19790606
                                          GB 1978-44319
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     GB 2008600
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     BR 7807431
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                           19790724
     NL 7811234
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                           19790517
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     FR 2408653
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     FR 2408653
                      В1
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     JP 54145290
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                           19791113
     JP 56028921
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     JP 57022322
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                                                            19801028
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     JP 57022323
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                      A2
                           19810711
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                                                            19801028
     JP 56085284
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ØS 4340678

Α

19820720

US 1981-222388

19810105

US 4352741 US 1981-269681 19821005 19810602 Α PRAI US 1977-851757 19771115 The displacement medium is a pyruvate-free xanthan gum [11138-66-2] slurry built up by aerobic fermn. of Xanthomonas campestris (ATCC 31313) in an aq. culture contg. a hydrocarbon and a source of N and trace elements until the slurry contains 100 ppm I. The slurry has a reduced ionic nature that can be further reduced by deacetylation, thus minimizing problems with Ca2+ in the underground formation. The slurry is used in injection solns. contg. .gtoreq.7% salt for displacement of oil from partially exhausted formations. 11138-66-2 TΤ RL: USES (Uses) (slurries, displacement media, for enhanced petroleum recovery) => fil wpix FILE 'WPIX' ENTERED AT 08:55:27 ON 10 OCT 2002 COPYRIGHT (C) 2002 THOMSON DERWENT FILE LAST UPDATED: 07 OCT 2002 <20021007/UP> 200264 MOST RECENT DERWENT UPDATE <200264/DW> DERWENT WORLD PATENTS INDEX SUBSCRIBER FILE, COVERS 1963 TO DATE >>> SLART (Simultaneous Left and Right Truncation) is now available in the /ABEX field. An additional search field /BIX is also provided which comprises both /BI and /ABEX <<< >>> The BATCH option for structure searches has been enabled in WPINDEX/WPIDS and WPIX <<< >>> PATENT IMAGES AVAILABLE FOR PRINT AND DISPLAY <<< >>> FOR DETAILS OF THE PATENTS COVERED IN CURRENT UPDATES, SEE http://www.derwent.com/dwpi/updates/dwpicov/index.html <<< >>> FOR A COPY OF THE DERWENT WORLD PATENTS INDEX STN USER GUIDE, PLEASE VISIT: http://www.stn-international.de/training center/patents/stn guide.pdf <<< >>> FOR INFORMATION ON ALL DERWENT WORLD PATENTS INDEX USER GUIDES, PLEASE VISIT: http://www.derwent.com/userguides/dwpi guide.html <<< => d all abeq tech abex tot L77 ANSWER 1 OF 8 WPIX (C) 2002 THOMSON DERWENT 1999-132209 [11] WPIX AN DNC C1999-038740 TТ Fluids for oil mining - comprises of a de-acetylated xanthane gum and a compound increasing the medium ionic strength. DC A11 A14 A97 E37 H01 Q49 ΙN LANGLOIS, B PΑ (RHOD) RHODIA CHIM CYC 82 A1 19990128 (199911)* EN C09K007-02 PΤ WO 9903948 33p RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW NL OA PT SD SE SZ UG ZW W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE

> GH GM HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US

UZ VN YU ZW

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A1 19990122 (199911)
                                                      C09K007-02
     FR 2766203
    AU 9887343 A 19990210 (199925)
NO 2000000208 A 20000317 (200025)
                                                      C09K007-02
                                                      C09K000-00
                   A1 20000510 (200027) FR
     EP 998540
                                                      C09K007-02
         R: AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE
     CN 1267320
                   A 20000920 (200063)
                                                      C09K007-02
    MX 2000000600 A1 20001001 (200158)
                                                      C08B037-00
ADT WO 9903948 A1 WO 1998-FR1514 19980710; FR 2766203 A1 FR 1997-9087
     19970717; AU 9887343 A AU 1998-87343 19980710; NO 2000000208 A WO
     1998-FR1514 19980710, NO 2000-208 20000114; EP 998540 A1 EP 1998-938729
     19980710, WO 1998-FR1514 19980710; CN 1267320 A CN 1998-808299 19980710;
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         E21B021-06; E21B043-22; E21B043-25
     ICS
AΒ
    WO
         9903948 A UPAB: 19990316
     Fluid free of guar for oil mining is claimed. It consists of de-
    acetylated xanthane gum, in a
    polypentamer form and a compound increasing the medium ionic
     strength. It further comprises a filtrate reducer and a fluidifying or
    dispersing agent with a concentration of 0-1 %, an oxygen sensor with a
     concentration of 0-0.25 %, all with respect to the total weight of the
     fluid.
          USE - As a filtrate for oil mining.
    Dwg.0/1
    CPI GMPI
FS
    AB; DCN
FΆ
    CPI: A10-E09; A12-W10A; E05-A; E05-B01; E31; E33; E34; H01-B06
MC
L77
    ANSWER 2 OF 8 WPIX (C) 2002 THOMSON DERWENT
    1998-120375 [11]
AN
                       WPTX
     1998-110557 [10]; 1998-120374 [11]
CR
    C1998-039555
DNC
     Composition of amorphous cellulose nano-fibrils - used as viscosity
TΤ
    modifier in food, cosmetic and detergent products and building materials,
    and in fluids used in oil extraction.
DC
    A11 A96 A97 D13 D21 D25 E19 H01 L02
    BENCHIMOL, J; CANTIANI, R; GUERIN, G; SENECHAL, A; VINCENT, I;
IN
    LANGLOIS, B
     (RHOD) RHODIA CHIM; (RHON) RHONE-POULENC CHIM; (BENC-I) BENCHIMOL J;
PA
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ADT WO 9802487 A1 WO 1997-FR1291 19970711; FR 2751659 A1 FR 1996-9062
   19960715; AU 9736974 A AU 1997-36974 19970711; EP 912634 A1 EP 1997-933723
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19970711, WO 1997-FR1291 19970711; SK 9900034 A3 WO 1997-FR1291 19970711, SK 1999-34 19970711; BR 9710338 A BR 1997-10338 19970711, WO 1997-FR1291 19970711; HU 9903102 A2 WO 1997-FR1291 19970711, HU 1999-3102 19970711; JP 2000503704 W WO 1997-FR1291 19970711, JP 1998-505682 19970711; AU 723465 B AU 1997-36974 19970711, WO 1997-FR1291 19970711; US 6224663 B1 WO 1997-FR1291 19970711, US 1999-214774 19990908; US 2001004869 A1 Div ex US 1999-214774 19990908, US 2001-782802 20010214; JP 3247391 B2 WO 1997-FR1291 19970711, JP 1998-505682 19970711 FDT AU 9736974 A Based on WO 9802487; EP 912634 A1 Based on WO 9802487; BR 9710338 A Based on WO 9802487; HU 9903102 A2 Based on WO 9802487; JP 2000503704 W Based on WO 9802487; AU 723465 B Previous Publ. AU 9736974, Based on WO 9802487; US 6224663 B1 Based on WO 9802487; US 2001004869 A1 Div ex US 6224663; JP 3247391 B2 Previous Publ. JP 200003704, Based on WO 9802487 PRAI FR 1996-11779 19960927; FR 1996-9062 19960715 ICM C08L001-00; C08L001-02; D21C009-00 A23L001-00; A23L001-03; A61K007-00; C04B024-00; C04B024-38; C09D101-02; C09K007-00; C11D001-68; C11D003-00; C11D003-22; C11D003-382 C08L001-02, C08L001:28; C08L001-02, C08L005:00; C08L001-02, C08L029:04; C08L001-02; C08L001:28; C08L001-02, C08L001:28; C08L001-02, C08L001:28; C08L001-02, C08L005:00; C08L001-02, C08L029:04 WO 9802487 A UPAB: 20010711 The composition comprises essentially amorphous cellulose nano-fibrils; at least one additive chosen from carboxylated cellulose having degree of substitution > 0.95, a natural polysaccharide, or a polyol; and optionally at least one co-additive. The content of additive and co-additive is at most 30 wt.% based on total weight of nano-fibrils, additive and any co-additive(s). Also claimed is a suspension of cellulose nano-fibrils obtained by dispersing the above composition. The suspension preferably has a rheo-fluidising type rheological profile. USE - Carboxylated cellulose and optionally co-additives, with amorphous cellulose nano-fibrils are used to preserve the rheo-fluidising rheological profile of a suspension of amorphous cellulose nano-fibrils having undergone a drying stage. The compositions and suspensions are used as additives in cosmetic, detergent and food formulations, in formulations for use in building, and in fluids used in oil extraction. Dwg.0/0 CPI AB; DCN CPI: A03-A04A; A12-R01; A12-V04; A12-W09; A12-W12A; A12-W12B; D03-H01; D03-H01J; D08-B; D11-B; E10-D03C; H01-B06; L02-D ANSWER 3 OF 8 WPIX (C) 2002 THOMSON DERWENT 1997-503082 [46] WPIX C1997-160017 Preparation of aqueous tar suspo-emulsion for cleaning tar/sludge comprises mixing viscous tar composition, inorganic solids and water and surface active agent and thickening water-soluble polymer. A14 A25 A97 D25 E19 H08 GUERIN, G; HILL, D P; LANGLOIS, B; PRUITT, T E; SANDERS, F L; PRUIT, T E; HILL, P D (RHON) RHONE-POULENC CHIM; (RHON) RHONE-POULENC INC; (RHOD) RHODIA CHIMIE; (RHOD) RHODIA INC; (RHOD) RHODIA CHIM 71 WO 9736970 A2 19971009 (199746)* EN 34p C10C000-00 RW: AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE HU IL IS JP KE KG KP KR KZ LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK TJ TM TR TT UA UG UZ VN AU 9738776 A 19971022 (199808) C10C001-00 NO 9803791 A 19981019 (199901) C10C001-00

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     19970220, HU 2000-245 19970220; AU 723738 B AU 1997-38776 19970220; MX
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     KR 1998-706461 19980819; US 6197837 B1 Provisional US 1996-11977P
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     2001-203174 19970220
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FDT
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     Based on WO 9736970; KR 99087075 A Based on WO 9736970; JP 2001527587 W
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     19960220; US 1999-263641
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IC
         C10G017-00; C10L001-00; C10L001-32
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         C10G017-10; C10G075-00; C23G001-02; C23G001-24
AΒ
          9736970 A UPAB: 19990224
    The preparation of an aqueous tar suspoemulsion involves mixing a mixture
     comprising: (a) a viscous tar composition of tar(s), inorganic solids and
     optionally water, (b) water and (c) at least one surface-active agent
     exhibiting an HLB of at least 10 and optionally at least one thickening
     water-soluble polymer with mol. wt. more than 10,000. The relative amounts
     of water, surface-active agent and polymer are such that the viscosity of
     the mixture is the same as or more than 0.1 of the viscosity of the tar. A
     process for fluidizing tars/sludges or cleaning tars/sludges from
     containers/vessels involves contacting the tar/sludge with an inorganic
     acid and a surfactant.
          USE - For use in fluidizing tars and sludges especially in cleaning
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ADVANTAGE - Residues can be conditioned in fluid form which can be

tars from reaction vessels, process equipment, transport container and

storage tanks.

diluted with water or with acid and which is stable on storage. Does not require any physical alteration to the tank to remove the tar/sludge. Tar/sludge can be easily recovered and the sulphuric acid can be regenerated. Dwg.0/0 FS CPI FA AB; DCN MC CPI: A03-C03; A07-B04; A08-S; A11-A03; A11-C; A12-W12B; D11-A12; D11-D01B; E05-G09C; E05-G09D; E07-D09C; E10-A03; E10-A09A; E10-A09B4; E10-A09B8; E10-A22; E10-B02D6; E10-B03B; E10-B04A2; E10-C04F; E10-D03C; E10-E04M1; E10-E04M3; H08-E05 ANSWER 4 OF 8 WPIX (C) 2002 THOMSON DERWENT L77 ΑN **1997-086653** [08] WPIX DNN N1997-071447 DNC C1997-028131 Viscous aq. particle transport fluid - comprising water, guar gum and a TΤ non-acetylated but otherwise unmodified xanthan heteropolysaccharide polymer.. DC A11 A97 D16 H01 Q49 ΙN HODGE, R M (DUPO) DU PONT DE NEMOURS & CO E I PACYC A 19970107 (199708)* PΙ US 5591699 9p E21B043-26 ADT US 5591699 A CIP of US 1993-21943 19930224, US 1994-360558 19941221 PRAI US 1994-360558 19941221; US 1993-21943 19930224 IC ICM E21B043-26 AR US 5591699 A UPAB: 19970220 A viscous aq. particle transport fluid comprises: a) water; b) 0.08-1.0 wt.% of guar gum, and c) 0.02-0.5 wt.% of a nonacetylated but otherwise unmodified xanthan heteropolysaccharide polymer of formula (I): M = H ion of an alkali metal ion and where the ratio of guar gum to the amt. of xanthan heteropolysaccharide polymer used is 2:1 to 5:1. USE - The transport fluid is used as a drilling fluid, a fracturing fluid or as a filter structure emplacement fluid in mines. ADVANTAGE - The xanthan heteropolysaccharide polymer imparts viscosity to the aq. particle transport fluid. Dwg.0/2 CPI GMPI FS FA AB; GI CPI: A03-A00A; A03-C02; A12-W10A; D05-C08; H01-B06A; H01-C03 MC L77 ANSWER 5 OF 8 WPIX (C) 2002 THOMSON DERWENT 1992-398877 [48] ΑN WPIX 1987-051527 [08]; 1987-291652 [41]; 1991-030738 [05] CR DNC C1992-177000 TΤ New modified xanthan gums - with non-natural acetylation and/or pyruvylation pattern or poly tetramer structure, produced by Xanthomonas campestris mutants. DC A11 A97 D16 D17 H01 DOHERTY, D H; FERBER, D M; HASSLER, R A; MARRELLI, J D; VANDERSLICE, R W; TN DOHERTY, D N (MONS) MONSANTO CO; (GETT-N) GETTY SCI DEV CO; (TEXC) TEXACO DEV CORP; PA (DOHE-I) DOHERTY D H; (FERB-I) FERBER D M; (HASS-I) HASSLER R A; (MARR-I) MARRELLI J D; (VAND-I) VANDERSLICE R W; (KELC) CP KELCO US INC CYC 21 A1 19921112 (199248)* EN PI WO 9219753 67p C12P019-06 RW: AT BE CH DE DK ES FR GB GR IT LU MC NL SE W: CA FI JP NO A 19931105 (199404) FI 9304902 C12P000-00 NO 9304044 A 19931108 (199408) C12P000-00 EP 584206 A1 19940302 (199409) EN C12P019-06 R: AT BE CH DE DK ES FR GB GR IT LI LU MC NL SE

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     1993-4044 19931108; EP 584206 A1 EP 1992-911566 19920501, WO 1992-US3448
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     1995-475823 19950607, US 2001-986803 20011113; CA 2108895 C CA
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FDT EP 584206 A1 Based on WO 9219753; JP 06507433 W Based on WO 9219753; FI
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PRAI US 1991-696732
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                                19850806; US 1995-475823
                                                           19950607; US
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    EP 211288; EP 410326; US 4296203; US 4713449; WO 8705939
REP
IC
     ICM A01N043-04; C07H013-02; C08B037-00; C12P000-00; C12P019-06
         C07G017-00; C07H001-00; C12N001-00; C12N001-20; C12P019-04
ICI
    C12P019-06, C12R001:64
         9219753 A UPAB: 20021001
AΒ
     (A) New water-soluble polysaccharides (I) comprise repeating
    pentamer units with a G:M:GA ratio of 2:2:1 (G = D-glucose, M = C
     D-mannose, GA = D-glucuronic acid), where the G gps. are linked in a
    beta-(1,4) configuration, the inner M gps. are linked in an alpha-(1,3)
     configuration primarily to alternate G gps., the GA gps. are linked in a
     beta-(1,2) configuration to the inner M gps., and the outer M gps. are
     linked to the GA gps. in a beta-(1,4) configuration.
          (B) New water-soluble polysaccharides (II) comprise repeating
     tetramer units with a G:M:GA ratio of 2:1:1, where the G gps. are linked
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(B) New water-soluble polysaccharides (II) comprise repeating tetramer units with a G:M:GA ratio of 2:1:1, where the G gps. are linked in a beta-(1,4) configuration, the M gps. are acetylated or not acetylated at the 6-0 position and are linked in an alpha-(1,3) configuration primarily to alternate G gps., and the GA gps. are linked in a beta-(1,3) configuration to the M gps.

USE/ADVANTAGE - The gums are useful as viscosifiers or thickeners, e.g. in foods, drilling fluids and enhanced oil recovery fluids. The non-pyruvylated gums give lower viscosities at low temps. than wild-type (acetylated/pyruvylated) gums while having comparable viscosities at high temps. The non-acetylated gums give higher viscosities than wild-type gums over a broad temp. range.

Dwg.0/12

CPI

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FA
     AB
MC
     CPI: A03-A; A09-A; D03-H01J; D06-H; H01-B06; H01-D06
ABEO US
          5514791 A UPAB: 19960618
     A water-soluble polysaccharide polymer comprising repeating
    pentamer units having a D-glucose:D-mannose:D-glucuronic acid
     ratio of about 2:2:1, wherein the D-glucose moieties are linked in a
     beta-[1,4] configuration, inner D-mannose moieties are linked in an
     alpha-[1,3] configuration primarily to alternate glucose moieties, the
     D-glucuronic acid moieties are linked in a beta-[1,2] configuration to
     said inner mannose moieties, and outer mannose moieties are linked to said
     glucuronic acid moieties in a beta-[1,4] configuration, wherein said inner
     mannose moieties are not acetylated and a portion of said outer mannose
     moieties are acetylated.
     Dwg.0/10
     ANSWER 6 OF 8 WPIX (C) 2002 THOMSON DERWENT
ΑN
     1987-291652 [41]
                        WPIX
CR
     1987-051527 [08]; 1991-030738 [05]; 1992-398877 [48]
DNC
    C1987-123874
ΤI
     New water soluble mutant forms of xanthan gum -
     partic. lacking acetate or pyruvate substitution, esp. useful in oil
     recovery processes.
DC
     All A97 Dl3 Dl6 Dl7 H01
ΙN
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PΑ
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CYC
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                                                      C12P019-06
                     19970729 (199742)
     CA 1339113
                   С
                                                      C12P019-06
     JP 2670256
                   B2 19971029 (199748)
                                               15p
                                                      C08B037-00
     NO 309102
                   B1 20001211 (200101)
                                                      C12P019-06
ADT
    WO 8705939 A WO 1987-US606 19870324; JP 63503198 W JP 1987-502340
     19870324; EP 323952 A EP 1987-902919 19870324; NO 9201787 A Div ex NO
     1987-4846 19870324, WO 1987-US606 19870324, NO 1992-1787 19920506; EP
     511690 A2 EP 1992-111024 19870324; NO 172945 B WO 1987-US606 19870324, NO
     1987-4846 19871120; EP 323952 B1 EP 1987-902919 19870324, WO 1987-US606
     19870324; DE 3787026 G DE 1987-3787026 19870324, EP 1987-902919 19870324,
     WO 1987-US606 19870324; EP 511690 A3 EP 1992-111024 19870324; JP 2559437
     B2 JP 1987-502340 19870324, WO 1987-US606 19870324; EP 765939 A2 Div ex EP
     1992-111024 19870324, EP 1996-119095 19870324; EP 765939 A3 Div ex EP
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1992-111024 19870324, EP 1996-119095 19870324; EP 511690 B1 Div ex EP

maier - 10 / 082555 1987-902919 19870324, EP 1992-111024 19870324, Related to EP 1996-119095 19870324; JP 09176205 A Div ex JP 1987-502340 19870324, JP 1996-70588 19870324; DE 3752096 G DE 1987-3752096 19870324, EP 1992-111024 19870324; CA 1339113 C CA 1987-532838 19870324; JP 2670256 B2 Div ex JP 1987-502340 19870324, JP 1996-70588 19870324; NO 309102 B1 WO 1987-US606 19870324, Div ex NO 1987-4846 19871120, NO 1992-1787 19920506 EP 511690 A2 Related to EP 323952; NO 172945 B Previous Publ. NO 8704846, Div in NO 9201787; EP 323952 B1 Based on WO 8705939; DE 3787026 G Based on EP 323952, Based on WO 8705939; JP 2559437 B2 Previous Publ. JP 63503198, Based on WO 8705939; DE 3752096 G Based on EP 511690; JP 2670256 B2 Previous Publ. JP 09176205; NO 309102 B1 Previous Publ. NO 9201787 PRAI US 1987-29090 19870323; US 1986-842945 19860324; US 1986-844435 19860326 6.Jnl.Ref; US 3000790; US 4296203; 11Jnl.Ref; EP 211288; WO 8705938; No-SR.Pub; 7.Jnl.Ref; 5.Jnl.Ref; EP 66961

REP

ICM C08B037-00; C12P019-06 IC

C07G017-00; C12N001-20; C12P001-04; C12P019-04

C12R001-64 ICA

FDT

C12R001:64; C12P019-06, C12R001:64; C12N001-20, C12R001:64; C12P019-06, ICI C12R001:64; C12N001-20, C12R001:64; C12P019-06, C12R001:64 AΒ

8705939 A UPAB: 20020823 A series of water-soluble polysaccharides contg. D-glucose (G), D-mannose (M) and opt. D-glucuronic acid (GA) are new. They have the following characteristics. (1) G: M:GA ratio 2:1:1; G linked beta-1,4; M linked alpha-1,3 (generally to alternate G) and GA linked beta-1,2 to M; (2) G:M:GA ratio 2:2:1; G linked beta-1,4; M, not acetylated at 6-0, linked alpha-1,3 (generally to alternate G); GA linked beta-1,2 to nonacetylated M, and a second M contg. a 4,6-ketal linked pyruvate gp. linked beta-1,4 to GA; (3) similar to (2) but first M is 6-O-acetylated and second M does not contain pyruvate; (4) similar to (3) but the second M does not contain pyruvate, and the polymer is at least 90% acetylated; (5) G:M ratio 2:1; G linked beta-1,4 and M (at least 90% being 6-0-acetylated) and linked alpha-1,3, generally to alternate G.

USE/ADVANTAGE - These gums are superior to normal xanthan gums in shear rate, salt tolerance and temp. variation of viscosity properties. They are esp. useful in sec. and tert. oil recovery, but can also be used as thickeners (in foods, cosmetics, medicines, paper sizes, drilling muds and printing inks), gelling agents and to reduce frictional drag of fluid in pipes.

FS CPI

FA AΒ

MC CPI: A03-A; A03-C02; A12-W10B; D03-H01J; D04-B03; H01-B06; H01-D09 ABEQ EP 323952 B UPAB: 19931118

A composition comprising a water-soluble polysaccharide polymer having a D-glucose; D-mannose; D-glucuronic acid ratio of about 2:2:1 wherein (1) the D-glucose moieties are linked in a beta-1,4)configuration, (2) the D-mannose moieties are linked in an alpha-(1,3)configuration generally to alternative glucose moieties, and (3) the D-glucuronic acid moieties are linked in a beta-(1,2)configuration to the mannose moieties. Dwg.0/3

ABEQ EP 511690 A UPAB: 19931202 A series of water-soluble polysaccharides contg. D-glucose (G), D-mannose (M) and opt. D-glucuronic acid (GA) are new. They have the following characteristics. (1) G: M:GA ratio 2:1:1; G linked beta-1,4; M linked alpha-1,3 (generally to alternate G) and GA linked beta-1,2 to M; (2) G:M:GA ratio 2:2:1; G linked beta-1,4; M, not acetylated at 6-0, linked alpha-1,3 (generally to alternate G); GA linked beta-1,2 to nonacetylated M, and a second M contg. a 4,6-ketal linked pyruvate gp. linked beta-1,4 to GA; (3) similar to (2) but first M is 6-O-acetylated and second M does not contain pyruvate; (4) similar to (3) but the second M does not contain pyruvate, and the polymer is at least 90% acetylated; (5) G:M ratio 2:1; G linked beta-1,4 and M (at least 90%

being 6-O-acetylated) and linked alpha-1,3, generally to alternate G.

USE/ADVANTAGE - These gums are superior to normal xanthan gums in shear rate, salt tolerance and temp. variation of viscosity properties. They are esp. useful in sec. and tert. oil recovery, but can also be used as thickeners (in foods, cosmetics, medicines, paper sizes, drilling muds and printing inks), gelling agents and to reduce frictional drag of fluid in pipes.

ABEQ EP 511690 B UPAB: 19970828
A process for preparing a water-soluble polysaccharide polymer comprising repeating pentamer units having a D-glucose: D-mannose:
D-glucuronic acid ratio of about 2:2:1, where the D-glucose moieties are linked in a beta- (1,4) configuration, inner D-mannose moieties are linked in an alpha- (1,3) configuration primarily to alternate glucose moieties, the D-glucuronic acid moieties are linked in a beta- (1,2) configuration to the inner mannose moieties, and outer mannose moieties are linked to the glucuronic acid moieties in a beta (1,4) configuration, where the polysaccharide polymer is not acetylated, the process comprising: (a) obtaining an acetylase deficient mutant of Xanthomonas; and (b) culturing the Xanthomonas under conditions sufficient to produce the polysaccharide polymer.

Dwg. 0/4

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Dwg.0/4
L77
    ANSWER 7 OF 8 WPIX (C) 2002 THOMSON DERWENT
ΑN
     1987-079865 [12]
                        WPIX
    C1987-033317
DNC
     Modification of microbial polysaccharide(s) esp. xanthan
TΙ
     gum - by adding nitric acid, heating, cooling and neutralising,
     giving saline-compatible thickener for oil recovery fluids.
DC
     All A97 D16 D17 H01 Q49
ΙN
     GROS, P; PIPON, R
     (RHON) RHONE POULENC SPECIALITES CHIM
PA
CYC
    18
                  A 19870219 (198712)*
PΙ
    AU 8661120
                                              34p
     FR 2586249
                   A 19870220 (198713)
     JP 62039643
                  A 19870220 (198713)
     EP 215692
                  Α
                     19870325 (198714)
                                              15p
        R: AT DE FR GB IT NL SE
                A 19870324 (198715)
     BR 8603842
     NO 8603244
                  A 19870309 (198716)
                  A 19870215 (198719)
     FI 8603290
     DK 8603850
                  A 19870215 (198720)
     CN 86105280
                  A 19870211 (198818)
                  A 19880401 (198919)
     ES 2000962
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R: AT DE FR GB IT NL SE

CA 1265791 A 19900213 (199014)

DE 3669531 G 19900419 (199017)

SU 1570650 A 19900607 (199107)

US 5010186 A 19910423 (199120)

A 19891010 (198950)

B 19900314 (199011)

ADT AU 8661120 A AU 1986-61120 19860813; FR 2586249 A FR 1985-12382 19850814; JP 62039643 A JP 1986-188797 19860813; EP 215692 A EP 1986-401699 19860730; ES 2000962 A ES 1986-50 19860813; US 4873323 A US 1986-896282 19860814; SU 1570650 A SU 1986-4027937 19860813; US 5010186 A US 1989-372819 19890913

FR

9p.

PRAI FR 1985-12382 19850814

US 4873323

EP 215692

REP 1.Jnl.Ref; EP 103483; EP 78621; FR 2551070; RO 84329; US 4299825 IC A61K031-71; C07G017-00; C08B005-02; C08B009-06; C08B037-00; C08L005-00; C10M145-40; C12N019-00; C12P019-06; E21B043-22

AB AU 8661120 A UPAB: 19930922
Prepn. of a polysaccharide (I) modified by heat treatment of an aq. compsn. contg. 0.05-35 wt.% (I) is characterised in that: (a) the compsn. is acidified with NHO3 to pH 2-0.1, pref. below 1.5; (b) the compsn. is heated to 50-100, pref. 60-90 deg.C for 5-60, pref. 10-45 min.; and (c)

the compsn. is cooled and base (e.g. NaOH, KOH or NH4OH) is added to pH 5-7. Aq. compsns. of the modified polysaccharides are claimed.

USE/ADVANTAGE - Useful for modifying xanthan gums and scleroglucans (claimed), either in whole fermentation broth or as a soln. of commercial (I). The treated prods. are useful for controlling the mobility for enhanced oil recovery, treatment improving the filterability and injectability of the aq. solns. without reducing viscosifiation ability and making them useful in average and low permeability formations. Treatment gives prods. which can be used in a saline medium without clogging.

FS CPI GMPI

FA AB

MC CPI: A03-A00A; A10-E; A12-W10B; D05-C08; D06-H; H01-B06

ABEQ EP 215692 B UPAB: 19930922

Process for the preparation of a high molecular weight polysaccharide of the homo- or heteropoly-saccharide type, obtained from the fermentation of a carbohydrate by the action of microorganisms, the said polysaccharide, characterised in that: (a) the composition is acidified by adding nitric acid until a pH of between 2 and 0.1 is obtained, (b) the composition is heated to a temperature of 50-100 deg. C for a period of between 5 and 60 minutes and (c) the composition is cooled and the pH is increased to a value of 5 to 7 by adding a base.

ABEQ US 4873323 A UPAB: 19930922

Prepn. of a modified polysaccharide bipolymer comprises (i) acidifying an aq. compsn. of a polysaccharide with nitric acid to pH 2-0.1 (ii) heat treating the acidified compsn. at 50-100 deg.C for 5-60 mins and (iii) cooling the resultant compsn. and adjusting the pH to 5-7 to give a modified polysacc--haride with a decreased no.of acetyl gps.

Pref. the aq. compsn. comprises 0.05-35 wt. % zo of polysaccharide and is a carbohydrate fermentation, broth or an aq. soln. of a powdered polysaccharide.

USE/ADVANTAGE - The deacetylated polysaccharides produced have improved viscosity, filterability and injectability and are useful for the recovery of oil from partially depleted oil deposits.

ABEQ US 5010186 A UPAB: 19930922

The prepn. of a modified polysaccharide biopolymer, comprises: (i) acidifying an aq. compsn. of a polysaccharide with nitric acid to pH 2-0.1; (ii) heat treating the acidified compsn. at 50-100 deg.C for 5-60 mins.; and (iii) cooling the heat-treated compsn. and adjusting the pH to 5-7.

The biopolymer pref. further comprises biocide and/or enzyme. USE - Subterranean oil deposits are recovered using the polysaccharide biopolymer.

L77 ANSWER 8 OF 8 WPIX (C) 2002 THOMSON DERWENT

AN 1979-22429B [12] WPIX

TI Aq. gels based on de acetylated xanthan and galactomannan - are stronger than those obtd. with acetylated xanthan, useful in foodstuffs, explosives, air-treating compsns. etc..

DC A11 D13 D17 D22 K04 P34

IN BRIGAND, G; KRAGEN, H

PA (CECA) CECA SA

CYC 6

PI EP 1192 A 19790321 (197912)*

R: CH DE GB NL

FR 2402678 A 19790511 (197924)

EP 1192 B 19811223 (198201) FR

R: CH DE GB NL

DE 2861455 G 19820211 (198207)

US 4369125 A 19830118 (198306)

PRAI FR 1977-27467 19770912

REP US 3384498; US 3677961 IC A23L001-04; A61L009-01; B01J013-00; C06B047-14; C08L005-00 AB 1192 A UPAB: 19930901 Prodn. of aq. gels based on galactomannan (I) and xanthan (II) uses (partially) deacetylated xanthan (IIa) and a (IIa): (I) ratio of 15:85 to 90:10. Pref. (I) is obtd. from carob, guar, tara and/or Espina corona gums. Carob gum when used pref. has a degree of polymsn. sufficient to give a viscosity of 20-6000 cP (1% soln. at 20 degrees C; Brookfield viscometer at 20 rpm). (IIa) may be obtd. as in US3000790. Gelling compsns. based on (I) and (IIa) are also claimed, as are gels obtd. as above contg. 0.1-4% of (I) + (IIa)). Prods. are stronger than previously possible using (II). They can be used in foodstuffs, e.g. in aspics or gelled desserts, and in explosive gels and air-treatment prods. FS CPI GMPI FΑ CPI: A03-A; A10-E09; A12-S; D03-H01J; D06-H; K04-A MC => fil dpci FILE 'DPCI' ENTERED AT 08:56:28 ON 10 OCT 2002 COPYRIGHT (C) 2002 THOMSON DERWENT FILE LAST UPDATED: 10 OCT 2002 <20021010/UP> MOST RECENT DERWENT DPCI UPDATE 200235 PATENTS CITATION INDEX, COVERS 1973 TO DATE CURRENT UPDATES CONTAIN CORRECTIONS TO PREVIOUSLY LOADED DOCUMENTS <<< >>> LEARNING FILE LDPCI AVAILABLE <<< => d all L78 ANSWER 1 OF 1 DPCI (C) 2002 THOMSON DERWENT AN 1999-132209 [11] DPCI DNC C1999-038740 TIFluids for oil mining - comprises of a de-acetylated xanthane gum and a compound increasing the medium ionic strength. DC All Al4 A97 E37 H01 Q49 ΙN LANGLOIS, B PA (RHOD) RHODIA CHIM CYC 82 PΤ A1 19990128 (199911)* EN WO 9903948 33p C09K007-02 RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW NL OA PT SD SE SZ UG ZW W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE GH GM HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZW FR 2766203 A1 19990122 (199911) C09K007-02 AU 9887343 A 19990210 (199925) C09K007-02 NO 2000000208 A 20000317 (200025) C09K000-00 EP 998540 A1 20000510 (200027) FR C09K007-02 R: AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE CN 1267320 A 20000920 (200063) C09K007-02 MX 2000000600 A1 20001001 (200158) C08B037-00 ADT WO 9903948 A1 WO 1998-FR1514 19980710; FR 2766203 A1 FR 1997-9087 19970717; AU 9887343 A AU 1998-87343 19980710; NO 2000000208 A WO 1998-FR1514 19980710, NO 2000-208 20000114; EP 998540 A1 EP 1998-938729 19980710, WO 1998-FR1514 19980710; CN 1267320 A CN 1998-808299 19980710;

MX 2000000600 A1 MX 2000-600 20000117

FDT AU 9887343 A Based on WO 9903948; EP 998540 A1 Based on WO 9903948

PRAI FR 1997-9087 19970717

IC ICM C08B037-00; C09K000-00; C09K007-02

ICS E21B021-06; E21B043-22; E21B043-25

FS CPI GMPI

CTCS CITATION COUNTERS

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PNC.DI	0	Cited Patents Count (by inventor)
PNC.DX	6 ·	Cited Patents Count (by examiner)
IAC.DI	0	Cited Issuing Authority Count (by inventor)
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CRC.X	1	Cited Literature References Count (by examiner)

CDP CITED PATENTS UPD: 20001030

Cited by Examiner

CITING PATENT	CA	r cited patent accno
EP 998540	 А	No Citations
WO 9903948	A Y	EP 765939 A 1987-291652/41
	PA:	(GETT-N) GETTY SCI DEV CO
	IN:	DOHERTY, D H; FERBER, D M; MARRELLI, J D; VANDERSLICE,
		R W; VANDERSLIC, R W
	Y	GB 1080248 A 1968-74899P/00
	PA:	(ESSO) ESSO PRODN RES CO
	Y	US 3096293 A
	IN:	JEANES
	Y	US 4186803 A 1980-12617C/07
	PA:	(TEXA-N) TEXAS BRINE CORP
	IN:	MONDSHINE, T C
	A	US 4218327 A 1980-64043C/36
	PA:	(SHEL) SHELL OIL CO
	IN:	WELLINGTON, S L
	Y	US 4868293 A 1987-051527/08
	PA:	(GETT-N) GETTY SCI DEV CO
	IN:	SHANON, P; VANDERSLICE, R W; VANDERSLIC, R W; SHANNON,
		p

REN LITERATURE CITATIONS UPR: 20001030

Citations by Examiner

CITING PATENT CAT CITED LITERATURE

EP 998540 A See references of WO 9903948A1

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L85
    ANSWER 1 OF 4 WPIX (C) 2002 THOMSON DERWENT
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    1987-051527 [08]
                       WPIX
     1987-291652 [41]; 1991-030738 [05]; 1992-398877 [48]
CR
DNN
    N1987-039074
                        DNC C1987-021433
TΙ
    New polysaccharide from Xanthomonas contg. no glucuronic acid - useful as
     viscosifier, esp. for oil recovery under high temp. or salt conditions.
DC
    A97 B04 D13 D16 D21 H01 Q49
     SHANON, P; VANDERSLICE, R W; VANDERSLIC, R W; SHANNON, P.
ΙN
PΑ
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    EP 211288
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    NO 8603135
                 A 19870302 (198715)
    DK 8603721
                  A 19870207 (198719)
                  A 19870207 (198719)
    FI 8603213
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                  A 19870417 (198721)
    US 4713449
                  A 19871215 (198806)
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    CA 1247033
                  A 19881220 (198904)
    US 4868293
                  A 19890919 (198947)
                                              12p
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                  A 19900123 (199008)
    CA 1264537
    CA 1279180
                   С
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                   B1 19921021 (199243)
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                  G 19921126 (199249)
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     FI 92719
                   B 19940915 (199437)
                                                     C12P019-06
     JP 2520881
                   B2 19960731 (199635)
                                              14p
                                                     C08B037-00
                                              12p
     JP 08239403
                   A 19960917 (199647)
                                                     C08B037-00
                                              11p
     JP 2746560
                   B2 19980506 (199823)
                                                     C12N001-20
ADT
    EP 211288 A EP 1986-109782 19860716; JP 62084102 A JP 1986-178978
     19860731; US 4713449 A US 1985-762878 19850806; US 4868293 A US 1987-99618
     19870922; US 5102561 A US 1991-715861 19910617; EP 211288 B1 EP
     1986-109782 19860716; DE 3686986 G DE 1986-3686986 19860716, EP
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1986-109782 19860716; FI 92719 B FI 1986-3213 19860806; JP 2520881 B2 JP 1986-178978 19860731; JP 08239403 A Div ex JP 1986-178978 19860731, JP

1995-341723 19860731; JP 2746560 B2 Div ex JP 1986-178978 19860731, JP 1995-341723 19860731

FDT DE 3686986 G Based on EP 211288; FI 92719 B Previous Publ. FI 8603213; JP 2520881 B2 Previous Publ. JP 62084102; JP 2746560 B2 Previous Publ. JP 08239403

PRAI US 1987-99618 19870922; US 1985-762878 19850806; US 1989-333285 19890405; US 1991-715861 19910617

REP 2.Jnl.Ref; A3...198739; No-Sr.Pub

IC ICM C12N001-20; C12P019-06

ICS C08L001-00; C12R001-64; E21B021-00; E21B043-22; G05D024-00

ICA C08B037-00; C09K003-00; C09K007-00; C12P019-04

ICI C12P019-04, C12R001:64; C12N001-20, C12R001:64; C12P019-04, C12R001:64; C12P019-04, C12R001:64

AB EP 211288 A UPAB: 20020823

A new, water-soluble polysaccharide polymer (I) contains no glucuronic acid (GA) and has D-glucose:D-mannose ratio about 2:1. The glucose is linked in thebeta-1,4-configuration and the mannose in the beta-1,3 configuration, generally to alternate glucose residues. Opt. the mannose gps. are 6-O-acetylated. Also new are microbiologically pure cultures of Xanthomonas able to produce (I), specifically the strains ATCC 53195 and 53196.% (I) is made by aerobic fermentation of a Xanthomonas strain, unable to incorporate GA, on a suitable growth medium, pref. at 28-32 deg.C and pH 6-7.5. (I) is recovered by pptn., e.g. using isopropanol or by ultrafiltration.

USE/ADVANTAGE - (I) is used to increase the viscosity of an aq. media, esp. in oil recovery processes, but also in foods, cosmetics, medical formulations, paper sizes, drilling muds, printing inks or to reduce frictional drag of fluid flow in pipes. Compared with xanthan gum, (I) is a better viscosifier, esp. at high temp. and/or under high salt conditions.

Dwg.0/5

FS CPI GMPI

FA AB

MC CPI: A03-A00A; A12-W10B; B04-B02B1; B04-C02; D05-C08; H01-B06; H01-D06 ABEQ EP 211288 B UPAB: 19930922

A water-soluble polysaccharide polymer containing essentially no glucuronic acid moieties having a D-glucose: D-mannose ratio of about 2:1, wherein the D-glucose moieties are linked in a beta-(1,4) configuration and the D-mannose moieties are generally linked to alternate glucose moieties in an alpha-(1,3) configuration.

1/5

ABEQ US 4713449 A UPAB: 19930922

Water-sol., polysaccharide polymer (I) comprises glucose and mannose moieties, having a D-glucose: D-mannose ratio of 2:1.

The D-glucose moieties are linked in a beta-(1,4)-configuration and the D-mannose moieties are linked in an alpha-(1,3) configuration, generally to alternate glucose moieties.

USE/ADVANTAGE - (I) is a better viscosifier of water than naturally occurring xanthan gum.

ABEQ US 4868293 A UPAB: 19930922

Prepn. of a water-soluble, polysaccharide polymer compsn. contg. no glucuronic acid gps., having D-glucose; D-mannose ratio 2:1, and having the D-glucose gps. linked in a beta-1,4 configuration and the D-mannose gps. in alpha-1,3 configuration, generally to alternate glucose moieties is claimed. A growth medium is inoculated with a genus Xanthomonas microorganism which cannot incorporate glucuronic acid into the polysaccharide polymer, and the medium is incubated at suitable temp. and dissolved oxygen content. Microbiologically pure culture of the microorganism is also claimed.

USE/ADVANTAGE - Polymer is esp. useful in mobility control solns. for enhanced oil recovery. It performs well at high temp. and at high salt levels. It can also be used in food, medicines, drilling muds, cosmetics, paper sizing, etc.

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ABEQ US 5102561 A UPAB: 19930922

A new process for increasing the viscosity of aq. media (foods, cosmetics, medicines, petroleum drilling fluids) comprises dissolving in it a polysaccharide polymer without gluocouronic acid moieties but having D-glucose:D-mannose ratio of about 2:1. The D-glucose moieties are linked in beta (1.4) configuration and the D-mannose moieties are generally linked to alternate glucose moieties in an alpha (1.3) configuration.

A new process for recovering oil from a subterranean formation comprises injecting a soln. of the above polysaccharide into a well to displace trapped oil from the porous rock and collecting it.

ADVANTAGE - The above polysaccharide is a more effective viscosifying agent than the conventionally used xanthan gum and also has wider salinity and temp. ranges. It may be obtd. by altering Xanthomonas mutants to change its biosynthetic pathway to produce the desired polymer instead of xanthan gum.

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=> d all abeg tech abex 2-4
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L85 ANSWER 2 OF 4 WPIX (C) 2002 THOMSON DERWENT
```

AN 1980-64043C [36] WPIX

TI Prepn. of viscosity-stabilised xanthan gum solns. - by using deoxygenated water and adding antioxidant and alcohol.

DC A11 A97 D16 D17 H01

IN WELLINGTON, S L

PA (SHEL) SHELL OIL CO

CYC 1

PI US 4218327 A 19800819 (198036)*

PRAI US 1976-673518 19760405; US 1978-903279 19780505

IC C09K003-00

AB US 4218327 A UPAB: 19930902

Prepn. of aq. solns. thickened with xanthan gum (XG) is carried out by (a) treating an aq. liq. to remove dissolved O2, (b) adding a water-soluble S-contg. antioxidant (I), (c) adding a water-soluble, readily oxidisable alcohol (II), and (d) adding XG.

Specifically, (I) and (II) are defined as being capable of protecting a soln. contg. XG, NaCl, a sulphite-type oxygen scavenger and 800 ppm of each of (I) and (II) from drastic (>25%) loss of viscosity when boiled at atmospheric pressure for 5 min.

The process is esp. applicable to XG solns. for use in oil recovery operations. The solns. retain practically constant viscosity when exposed to high temps. over long periods.

FS CPI

FA AB

MC CPI: A03-A; A12-W10; D04-B03; D06-H; H01-D06

L85 ANSWER 3 OF 4 WPIX (C) 2002 THOMSON DERWENT

AN 1980-12617C [07] WPIX

CR 1979-87479B [48]

TI Well completion and workover method - using treating fluid comprising satd. aq. saline soln. with sized particles of water-soluble salt(s).

DC H01 Q49

IN MONDSHINE, T C

PA (TEXA-N) TEXAS BRINE CORP

CYC :

PI US 4186803 A 19800205 (198007)*

PRAI US 1976-735169 19761026; US 1977-850639 19771111; US 1978-938033 19780830

IC E21B033-13

AB US 4186803 A UPAB: 19930902

In a well completion and workover method in which a subterranean formation in a well is contacted with a treating fluid, (a) a treating fluid comprising a satd. aq. saline soln. with >=1 water-soluble salt which is

insoluble in the saline soln. is pumped in the well so that it contacts the formation, the saline soln. and the salt each being selected from KCl, NaCl, CaCl2, Na2SO4, Na2CO3, NaHCO3, CaBr2 and K2CO3 and mixts. of these; (b) the salt is maintained in a particle size range of 5-800 mu with >5% of the particles being coarser than 44 mu to bridge and seal off the formation; and (c) the salt bridging particles are dissolved off the formation to remove the bridge and seal from the formation for flow of hydrocarbons into the well.

Pref. a viscosifier and suspension additive (e.g. CMC) in an amt. of 0.2--5 lb. per bbl. of satd. brine soln., and a fluid loss additive are also circulated in the well bore with the treating fluid. US4175042 claimed the treating fluid.

FS CPI GMPI

FA AB

MC CPI: H01-C; H01-C10

L85 ANSWER 4 OF 4 WPIX (C) 2002 THOMSON DERWENT

AN 1968-74899P [00] WPIX

TI Aqueous medium for use in underground boring operations.

DC A97 D16 H01

PA (ESSO) ESSO PRODN RES CO

CYC 2

PI NL 6608742 A (196800)*
GB 1080248 A (196801)
NL 148083 B 19751225 (197608)

<--

PRAI US 1965-466207 19650623

IC C08B037-00; C08L005-00; C09K007-02

AB NL 6608742 A UPAB: 19930831

Process for injecting an aqueous medium into a borehole in the earth, so that the aqueous medium contacts an underground formation, employs an aqueous medium containing a heteropolysaccharide, produced by the action of bacteria of the genus Xanthomonas on a carbohydrate. The heteropolysaccharide is crosslinked in the medium with polyvalent cations of a metal of Groups III-VIII.

The heteropolysaccharide is readily produced by fermentation of X. Campestris upon a carbohydrate containing aqueous medium, under conventional aerabic conditions. The product may be obtained from the filtered liquor by means of precipitation with methanol or ethanol. The heteropolysaccharide is obtained as a voluminous powder.

FS CPI

FA AB

MC CPI: A03-A; A11-C02; A12-A02

=> fil hcaplus

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FILE COVERS 1907 - 10 Oct 2002 VOL 137 ISS 15 FILE LAST UPDATED: 9 Oct 2002 (20021009/ED)

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=> d all 186

L86 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2002 ACS

AN 1963:436371 HCAPLUS

DN 59:36371

OREF 59:6604a

TI Deacetylated polysaccharide thickeners

IN Jeanes, Allene R.; Sloneker, James H.

PA United States Dept. of Agriculture

SO 3 pp.; Division of U.S. 3,000,790 (CA 56, 2625d)

DT Patent

LA Unavailable

NCL 252316000

CC 50 (Industrial Carbohydrates)

PATENT NO. KIND DATE APPLICATION NO. DATE

PI US 3096293 19630702 US 19601031 <--

AB The disclosures are similar, but the claims are different.

=> fil tulsa

FILE 'TULSA' ENTERED AT 09:05:45 ON 10 OCT 2002 COPYRIGHT (C) 2002 The University of Tulsa (UTULSA)

FILE COVERS 1965 TO 9 OCT 2002 (20021009/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d bib ab rn tot

L94 ANSWER 1 OF 9 TULSA COPYRIGHT 2002 UTULSA

AN 2001:11565 TULSA

DN 753893

TI NOVEL APPLICATION OF SYNERGISTIC GUAR/NON- ACETYLATED XANTHAN GUM MIXTURES IN HYDRAULIC FRACTURING

AU FISCHER, C C; CONSTIEN, V G; NAVARRETE, R C; COFFEY, M D; ASADI, M

CS CONSTIEN & ASSOCIATES; KELCO OIL FIELD GROUP; CP KELCO; STIM LAB INC

SO SPE OILFIELD CHEM INT SYMP (HOUSTON, TX, 2/13-16/2001) PROC 2001 (SPE-65037; AVAILABLE ON CD-ROM; COLOR; 12 PP; 14 REFS)

DT Conference; Conference Article

LA English

AB Fracturing fluids have traditionally been viscosified with guar and guar derivatives. Non-acetylated xanthan is a variant of xanthan gum which when combined with guar in solution develops a synergistic interaction that generates superior viscosity and particle transport at lower polymer concentrations. These water-base linear fluids have improved low shear viscosity at concentrations at or below 25 lb/1,000 gal when compared to fluids viscosified using a single viscosifier such as guar or xanthan gum. The polymer mixtures can be crosslinked to provide enhanced

viscosity at higher temperatures. The effect of different parameters on the rheology of the mixtures is presented, such as the ratio of guar to non-acetylated xanthan, the effect of salts, temperature and shear history. Large scale proppant transport tests were performed to evaluate proppant transport of the mixtures with respect to pure guar fluids RN 9000-30-0 (GUAR GUM) (GALACTOMANNAN) 11078-30-1 11138-66-2 (XANTHAN GUM) L94 ANSWER 2 OF 9 TULSA COPYRIGHT 2002 UTULSA 1999:22484 TULSA ΑN 711685 DN CR 704454 TΤ (R) FLUIDS USEFUL FOR THE EXPLOITATION OF PETROLEUM COMPRISING DE -ACETYLATED XANTHANE GUM AND AT LEAST ONE COMPOUND INCREASING THE IONIC STRENGTH OF THE MEDIUM (FLUIDES UTILISABLES DANS L'EXPLOITATION DU PETROLE COMPRENANT DE LA GOMME XANTHANE DESACETYLEE ET AU MOINS UN COMPOSE AUGMENTANT LA FORCE IONIQUE DU MILIEU) IN LANGLOIS, B PARHODIA CHIMIE PΙ FR 2766203 19990122 ΑI FR 1997-9709087 19970717 FR 2,766,203, C 1/22/1999, F 7/17/1997 (APPL 9,709,087) (C09K-007/02; SO E21B-021/06; E21B-043/25; E21B-043/22) BULL OFFIC PROPRIETE IND (FR) NO 3, P 88, 1/22/1999 (ISSN 07507650; IN FRENCH; ABSTRACT ONLY) (AO) SRPA# 704,454 DTPatent LA French AΒ (For abstract and indexing, see Abstract #704,454) ANSWER 3 OF 9 TULSA COPYRIGHT 2002 UTULSA L94 ΑN 1999:15253 TULSA 704454 DN FLUIDS USEFUL FOR OIL MINING COMPRISING DE- ACETYLATED TΙ XANTHANE GUM AND AT LEAST ONE COMPOUND INCREASING THE MEDIUM IONIC STRENGTH (FLUIDES UTILISABLES DANS L'EXPLOITATION DU PETROLE COMPRENANT DE LA GOMME XANTHANE DESACETYLEE ET AU MOINS UN COMPOSE AUGMENTANT LA FORCE IONIQUE DU MILIEU) IN LANGLOIS, B PA RHODIA CHIMIE PI WO 9903948 19990128 19980710 AΙ PRAI FR 1997-9709087 19970717 WORLD 99/03,948, P 1/28/1999, F 7/10/1998, PR FR 7/17/1997 (APPL SO 9,709,087) (C09K-007/02; C08B-037/00) (33 PP; 21 CLAIMS; IN FRENCH) DΤ Patent LAFrench AB Guar-free fluids capable of being used in oil exploitation are described that contain deacetylated xanthane gum in the form of a polypentamer, combined with at least one compound increasing the medium ionic strength. The fluids can be used as filtrate reducers with enhanced properties, using deacetylated xanthane

completion fluids.
RN 11138-66-2 (XANTHAN)
11138-66-2 (XANTHAN GUM)

L94 ANSWER 4 OF 9 TULSA COPYRIGHT 2002 UTULSA

qum in the form of a polypentamer, combined with a compound

increasing the medium ionic strength and a standard filtrate reducer. The fluids have application as an additive in drilling fluids, workover and

AN 97:15758 TULSA

DN 655592

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CR
     545426
     (R) FAMILY OF XANTHAN-BASED POLYSACCHARIDE POLYMERS INCLUDING
ΤI
    NON-ACETYLATED AND/OR NON-PYRUVYLATED GUM
IN
     DOHERTY, D H; FERBER, D M; MARRELLI, J D
     GETTY SCIENTIFIC DEV CO
PΑ
     EP 765939 19970402
PΤ
     ΕP
ΑT
               19870324
PRAI US 1986-842945 19860324
PRAI US 1986-844435 19860326
PRAI US 1987-29090 19870323
     EUROPE 765,939, P 4/2/97, F 3/24/87, PR US 3/24/86 (APPL 842,945), US
     3/26/86 (APPL 844,435) AND US 3/23/87 (APPL 29,090) (C12P-019/06;
     C08B-037/00; C12R-001/64) EUROPE PAT BULL V 1997, NO 14, P 94, 4/2/97
     (ISSN 01709305; ABSTRACT ONLY) (AO) SRPA# 545,426
DΤ
     Patent
LA
     English
AB ·
    (For abstract and indexing, see Abstract #545,426)
RN
     11138-66-2 (XANTHAN GUM)
    ANSWER 5 OF 9 TULSA COPYRIGHT 2002 UTULSA
L94
ΑN
     97:5445 TULSA
DN
     645279
     PARTICLE TRANSPORT FLUIDS THICKENED WITH ACETYLATE FREE XANTHAN
TΙ
     HETEROPOLYSACCHARIDE BIOPOLYMER PLUS GUAR GUM
IN
     HODGE, R M
PA
     DU PONT DE NEMOURS & CO
PΙ
     US 5591699 19970107
     US
AΙ
               19941221
PRAI US 1993-21943 19930224
     US 5,591,699, C 1/7/97, F 12/21/94, PR US 2/24/93 (APPL 21,943)
SO
     (F21B-043/26) (9 PP; 8 CLAIMS)
DT
     Patent
LA
    English
AB
    A quite small amount of nonacetylated but otherwise unmodified
     xanthan heteropolysaccharide polymer plus a quite small amount of
     guar gum has been found to impart viscosity to an aqueous particle
     transport fluid (for example, a drilling fluid, fracturing fluid, or a
     filter structure emplacement fluid) sufficient to suspend mineral
     particles. A crosslinking agent can also be employed to further decrease
     the amount of non-acetylated xanthan,
    heteropolysaccharide polymer and guar gum that are needed for particle
     suspension. These xanthan polymers can be produced by a
    Xanthomonas campestris variant such as that having Accession No. 68038 at
     the American Type Culture Collection, Rockville, Md., or by a mutant X.
     campestris variant having a suitable chromosomal deletion mutation. The
     xanthan polymer variants which are suitable are quite specific.
     For example, xanthan polymers produced by native or wild X.
     campestris are nowhere nearly as effective in suspending particles, nor
     are other variants produced by genetic engineering means, nor are
     conventional xantham polymers which are deacetylated by chemical
    means.
     9000-30-0 (GUAR GUM)
RN
       11138-66-2
                   (XANTHAN)
       11138-66-2
                   (XANTHAN GUM)
     39421-75-5 (HYDROXYPROPYL GUAR GUM)
L94
    ANSWER 6 OF 9 TULSA COPYRIGHT 2002 UTULSA
     93:5013 TULSA
AN
DN
     545426
     FAMILY OF XANTHAN-BASED POLYSACCHARIDE POLYMERS INCLUDING
TΙ
    NON-ACETYLATED AND/OR NON-PYRUVYLATED GUM
     DOHERTY, D H; FERBER, D M; MARRELLI, J D; VANDERSLICE, R W
ΤN
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PΑ

GETTY SCIENTIFIC DEV CO

```
EP 511690 19921104
PΙ
ΑI
    EΡ
               19870324
PRAI US 1986-842945 19860324
PRAI US 1986-844435 19860326
PRAI US 1987-29090 19870323
     EUROPE 511,690, P 11/4/92, F 3/24/87, PR US 3/24/86 (APPL 842,945), US
     3/26/86 (APPL 844,435) AND US 3/23/87 (APPL 29,090) (C12P-019/06;
     C12R-001/64) (17 PP; 16 CLAIMS)
DT
     Patent
LA
     English
AB
     Variant xanthan gums are described, including
     non- acetylated, non-pyruvylated, non-
     acetylated and non- pyruvylated, and fully-acetylated
     xanthan gums. In addition, in vitro and in vivo methods
     for the synthesis of these gums are described. Mutant Xanthomonas
     campestris strains useful in these syntheses are also specified. The gums
     are commonly used as thickening agents, in secondary or tertiary oil
     recovery as a mobility control and profile modification agent, and in
     drilling fluids.
RN
     127-17-3 (PYRUVIC ACID)
     7782-44-7
                (OXYGEN)
       11138-66-2 (XANTHAN GUM)
     25777-71-3 (NATURAL RESIN)
     50-99-7Q, 25191-16-6Q (GLUCOSE)
     3458-28-4Q, 31103-86-3Q
                             (MANNOSE)
L94
    ANSWER 7 OF 9 TULSA COPYRIGHT 2002 UTULSA
ΑN
     87:10103 TULSA
DN
     420704
     A POLYSACCHARIDE POLYMER MADE BY XANTHOMONAS
ΤI
IN
     VANDERSLICE, R W; SHANON, P
PA
     GETTY SCIENTIFIC DEV CO
ΡI
     EP 211288 19870225
ΑI
     EΡ
               19860716
PRAI US 1985-762878 19850806
     EUROP 211,288, P 2/25/87, F 7/16/86, PR US 8/6/85 (APPL 762,878) (29 PP;
     18 CLAIMS)
DΤ
     Patent
LA
     English
AB
    A polysaccharide polymer, which is a viscosifier of water, and its
    non-acetylated form are comprised of glucose and mannose
    moieties in a ratio of 2:1. Also described are Xanthomonas mutants which
     produce the polysaccharide polymer but which do not produce
     xanthan gum. A method of preparing the polysaccharide
     polymers is described. The polymers may be used in enhanced oil recovery
     as mobility control and profile modification agents and in drilling
     fluids.
L94
    ANSWER 8 OF 9 TULSA COPYRIGHT 2002 UTULSA
AΝ
     82:8280 TULSA
DN
     318268
     XANTHOMONAS BIPOLYMER FOR USE IN DISPLACEMENT OF OIL FROM PARTIALLY
TΤ
     DEPLETED RESERVOIRS
IN
     WERNAU, W C
PΙ
     US 4296203 19811020
ΑI
     US 1977-851757 19771115
     US 4,296,203, C 10/20/81, F 11/15/77 (APPL 851,757) (PFIZER INC) (9
SO
     CLAIMS)
DT
     Patent
LA
     English
AB
     A process is described for preparing a pyruvate-free Xanthomonas
     colloid-containing fermentation broth suitable for the preparation of
```

mobility control solutions used in oil recovery which comprises

aerobically fermenting a mutant strain of the genus Xanthomonas in an aqueous nutrient medium. The pyruvate-free xanthan and the deacetylated form of this biopolymer provide mobility control solutions which are especially useful for enhanced oil recovery where high brine applications are involved. The mobility control solutions produced in accord with this process are employed in oil recovery in the same manner as previously known mobility control solutions. (9 claims) 127-17-3 (PYRUVIC ACID) RN (XANTHAN GUM) 11138-66-2 25777-71-3 (NATURAL RESIN) ANSWER 9 OF 9 TULSA COPYRIGHT 2002 UTULSA L94 ΑN 82:7340 TULSA DN 317328 ΤI XANTHOMONAS BIOPOLYMER FOR USE IN DISPLACEMENT OF OIL FROM PARTIALLY DEPLETED RESERVOIRS IN WERNAU, W C CA 1113875 19811208 PΙ 19781024 ΑI PRAI US 1977-851757 19771115 CAN 1,113,875, C 12/8/81, F 10/24/78, PR US 11/15/77 (APPL 851,757) SO (PFIZER INC) (16 CLAIMS) DΤ Patent English LA A process is described for preparing a pyruvate-free Xanthomonas AB colloid-containing fermentation broth suitable for the preparation of mobility control solutions used in secondary oil recovery. The process involves aerobically fermenting a pyruvate-free xanthan -producing strain of a species of the genus Xanthomonas in an aqueous nutrient medium. The ingredients of the medium comprise a carbohydrate, a source of assimilable nitrogen, and trace elements. The fermentation is continued until at least 100 ppm of pyruvate-free xanthan is present in the broth, and where required deacetylation is performed. The process also provides a pyruvate-free xanthan -containing fermentation broth, pyruvate-free xanthan, and their deacetylated forms. The pyruvate-free xanthan and the deacetylated form of this biopolymer provide mobility control solutions which are especially useful for enhanced oil recovery where high brine applications are involved. (16 claims) 127-17-3 (PYRUVIC ACID) RN 8002-05-9 (CRUDE OIL) 8002-05-9 (PETROLEUM) 11138-66-2 (XANTHAN GUM) 25777-71-3 (NATURAL RESIN) => d his (FILE 'HOME' ENTERED AT 07:45:35 ON 10 OCT 2002) SET COST OFF FILE 'REGISTRY' ENTERED AT 07:46:08 ON 10 OCT 2002 E XANTHAN GUM/CN L1 1 S E3 FILE 'HCAPLUS' ENTERED AT 07:46:23 ON 10 OCT 2002 6898 S L1 L2 L3 20 S L1 (L) DEACET? 20 S L3 AND L2 1.4 L56651 S XANTHAN (A) GUM E DEACETYLATION/CT E E3+ALL L6 1069 S E5

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E E4+ALL
L7
            597 S E4
                 E E9+ALL
L8
            987 S E4, E3+NT
                 E E10+ALL
                 E E8+ALL
            1466 S E2, E4
L9
L10
             14 S L2, L5 AND L6-L9
             43 S L2, L5 AND DEACET?
L11
L12
             13 S L2, L5 AND DEACYL?
L13
              57 S L4, L10-L12 AND L2-L12
                 E LANGLOIS B/AU
              67 S E3-E5, E11-E13
L14
L15
              1 S L14 AND L13
L16
              1 S L14 AND L2, L5
                 E RHODIA/PA,CS
           1137 S E3, E4
L17
             32 S L2, L5 AND L17
L18
              1 S L13 AND L15, L16, L18
L19
L20
             31 S L18 NOT L19
                 E DRILLING FLUID/CT
                 E E4+ALL
L21
           7554 S E2, E3, E1+NT
                 E E8+ALL
L22
           1151 S E2, E1+NT
                 E E7+ALL
L23
           1561 S E3+NT
                 E DRILLING FLUID/CT
                 E E4+ALL
                 E E9+ALL
L24
           2374 S E4, E3+NT
                 E E13+ALL
L25
           5086 S E4, E3+NT
                 E E12+ALL
L26
           2374 S E4, E3+NT
                 E E2+ALL
L27
          .12718 S E3, E2+NT
                 E DRILLING FLUIDS/CT
L28
            402 S E8
                 E E3+ALL
                 E E11+ALL
L29
            388 S E1
            276 S L2, L5 AND L21-L29
L30
              3 S L30 AND L20
L31
              1 S L30 AND L13
L32
L33
              4 S L31, L32
L34
              1 S L33 AND LANGLOIS ?/AU
             56 S L13 NOT L34
L35
L36
              0 S L35 AND L30
              1 S L35 AND FUEL?/SC,SX
L37
              2 S L34, L37
L38
L39
             55 S L35 NOT L38
L40
             55 S L39 AND XANTHAN
             47 S L40 AND GUM
L41
              8 S L40 NOT L41
L42
L43
             55 S L39-L42
              49 S L43 AND (PD<=20000114 OR PRD<=20000114 OR AD<=20000114)
L44
                 SEL DN AN 3 4 14 25 30 31 35 44 45 46 47
L45
             11 S L44 AND E1-E33
L46
             13 S L38, L45
L47
              44 S L43 NOT L46
                 SEL DN AN 5 6 11 16 17 18 19 20 25 26 28 31 32 33 34 35 38 39 4
L48
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L49
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L51
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              5 S L50 NOT L51
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              4 S L52 NOT SUBSTRATE/TI
L54
             38 S L49, L51, L53
L55
           8536 S L2, L5 OR XANTHAN
             3 S L55 AND ?PENTAMER?
L56
L57
             40 S L54, L56
             40 S L57 AND L2-L57
L58
              9 S L58 AND ?ACYL?
L59
L60
             36 S L58 AND ?ACETYL?
L61
             40 S L58-L60
            161 S L55 AND C09K007/IC, ICM, ICS
L62
L63
              1 S L62 AND L61
             40 S L61, L63
L64
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     FILE 'WPIX' ENTERED AT 08:21:47 ON 10 OCT 2002
                E LANGLOIS B/AU
L65
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L66
           4808 S XANTHAN? (A) GUM OR ?XANTHAN? OR R16377/DCN, PLE
L67
              3 S L65 AND L66
                E R07345+ALL/DCN
                E R90071+ALL/DCN
                E R90015+ALL/DCN
                E R90000+ALL/DCN
                E R90083+ALL/DCN
                E R12062+ALL/DCN
L68
             38 S (R16377(L)M2095)/PLE
L69
            219 S (R16377(L)M2391)/PLE
L70
             26 S L66 AND (DEACETY? OR DE ACETY? OR NONACETY?) OR NON ACETY?)
L71
              3 S L66 AND ?PENTAMER?
L72
              1 S L68, L69 AND L70, L71
L73
              3 S L67, L72
L74
             37 S L68 NOT L73
L75
             25 S L70 NOT L73, L74
                SEL DN AN 13 14 18 20 25
L76
              5 S L75 AND E1-E10
L77
              8 S L73, L76 AND L65-L76
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              1 S E3
L78
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     FILE 'WPIX' ENTERED AT 08:57:00 ON 10 OCT 2002
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L79
              1 S E3
                E EP765939/PN
L80
              1 S E3
                E GB1080248/PN
L81
              1 S E3
                E US3096293/PN
                E US4186803/PN
L82
              1 S E3
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E US4218327/PN

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L83
              1 S E3
                E US4868293/PN
L84
              1 S E3
L85
              4 S L79-L84 NOT L77
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     FILE 'HCAPLUS' ENTERED AT 09:00:09 ON 10 OCT 2002
                E US3096293/PN
              1 S E3
L86
L87
              1 S L86 NOT L64
L88
              0 S L87 AND L2-L64
     FILE 'HCAPLUS' ENTERED AT 09:00:55 ON 10 OCT 2002
     FILE 'TULSA' ENTERED AT 09:03:10 ON 10 OCT 2002
L89
            994 S L1
           1051 S XANTHAN? (A) GUM OR XANTHAN?
L90
L91
           1051 S L89, L90
                E XANTHAN/CT
                E E3+ALL
            994 S E7, E8
L92
           1051 S L91, L92
L93
L94
              9 S L93 AND (DEACET? OR NONACET? OR (DE OR NON) () ACET?)
                E DEACET/CT
                E ACET/CT
     FILE 'TULSA' ENTERED AT 09:05:45 ON 10 OCT 2002
     INDEX '1MOBILITY, 2MOBILITY, ADISALERTS, AEROSPACE, AGRICOLA, ALUMINIUM,
     ANABSTR, AQUASCI, AQUIRE, BABS, BIBLIODATA, BIOBUSINESS, BIOCOMMERCE,
     BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, BLLDB, CABA, CANCERLIT, CAOLD,
     CAPLUS, CASREACT, CBNB, CEABA-VTB, ... 'ENTERED AT 09:08:24 ON 10 OCT 2002
     FILE 'CBNB, CEN' ENTERED AT 09:08:45 ON 10 OCT 2002
            151 S L1
L95
L96
            251 S XANTHAN? (A) GUM OR XANTHAN?
L97
            251 S L96, L95
              O S L97 AND (DEACET? OR NONACET? OR (DE OR NON) () ACET?)
L98
              O S L97 AND (DEACYL? OR NONACYL? OR (DE OR NON) () ACYL?)
L99
              O S L97 AND (PENTAMER? OR POLYPENTAMER?)
L100
L101
              6 S L97 AND ACET?
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